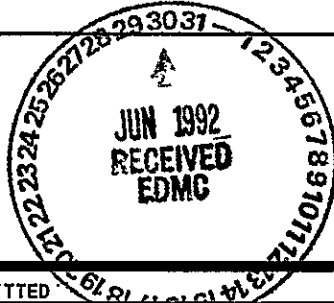


 START STA-4 APR 20 1992	ENGINEERING DATA TRANSMITTAL	0021622 Page 1 of 1 1. EDT 157741
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2. To: (Receiving Organization) Distribution	3. From: (Originating Organization) 200 Area Steam & Water Utilities	4. Related EDT No.: 150632
5. Proj./Prog./Dept./Div.: WHC-SD-WM-PLN-034	6. Cog. Engr.: D.R. Herman	7. Purchase Order No.: N/A
8. Originator Remarks: Approval and release of the Sampling and Analysis Plan for 284E Area Power Plant.		9. Equip./Component No.: N/A
		10. System/Bldg./Facility: N/A
11. Receiver Remarks:		12. Major Assm. Dwg. No.: N/A
		13. Permit/Permit Application No.: N/A
		14. Required Response Date: N/A




15. DATA TRANSMITTED					(F)	(G)	(H)	(I)
(A) Item No.	(B) Document/Drawing No.	(C) Sheet No.	(D) Rev. No.	(E) Title or Description of Data Transmitted	Impact Level	Reason for Transmittal	Originator Disposition	Receiver Disposition
1	WHC-SD-WM-PLN-034		Final Draft Rev 0	SAP for 284E Area Power Plant	3	1,2		

16. KEY											
Impact Level (F)			Reason for Transmittal (G)				Disposition (H) & (I)				
1, 2, 3, or 4 (see MRP 5.43)			1. Approval 2. Release 3. Information 4. Review 5. Post-Review 6. Dist. (Receipt Acknow. Required)				1. Approved 2. Approved w/comment 3. Disapproved w/comment 4. Reviewed no/comment 5. Reviewed w/comment 6. Receipt acknowledged				
(G)	(H)	17. SIGNATURE/DISTRIBUTION (See Impact Level for required signatures)								(G)	(H)
Reason	Disp.	(J) Name	(K) Signature	(L) Date	(M) MSIN	(J) Name	(K) Signature	(L) Date	(M) MSIN	Reason	Disp.
1	1	SA Brisbin	<i>SA Brisbin</i>	4/17/92	H4-16	RW Oldham	<i>RW Oldham</i>	4/16/92	H4-57	1	2
1	1	DR Herman	<i>DR Herman</i>	4/16/92	S4-01	EF Riedel	<i>EF Riedel</i>	4/16/92	S4-53	1	1
1	1	DM Nguyen	<i>DM Nguyen</i>	4/16/92	R1-48	E Yulis	<i>E Yulis</i>	4/16/92	S4-01	1	1
1	1	WA White	<i>WA White</i>	4/16/92	A7-27						

18. Signature of EDT Originator <i>DR Herman</i> 4/16/92	19. Authorized Representative for Receiving Organization _____ Date	20. Cognizant/Project Engineer's Manager <i>E Yulis</i> 4-16-92	21. DOE APPROVAL (if required) Ltr. No.
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			ID Number (include revision, volume, etc.) WHC-SD-WM-PLN-034 Rev 0		
			List attachments. N/A		
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Title Sampling and Analysis Plan of the 284E Powerplant Shop Process Wastewater Streams		Unclassified Category UC-		Impact Level 3	
New or novel (patentable) subject matter? <input checked="" type="checkbox"/> No <input type="checkbox"/> Yes If "Yes", has disclosure been submitted by WHC or other company? <input type="checkbox"/> No <input type="checkbox"/> Yes Disclosure No(s).		Information received from others in confidence, such as proprietary data, trade secrets, and/or inventions? <input checked="" type="checkbox"/> No <input type="checkbox"/> Yes (Identify)			
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Date(s) of Conference or Meeting		City/State		Will proceedings be published? <input type="checkbox"/> Yes <input type="checkbox"/> No Will material be handed out? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Title of Journal					
CHECKLIST FOR SIGNATORIES					
Review Required per WHC-CM-3-4		Reviewer - Signature Indicates Approval			
	Yes	No	Name (printed)	Signature	Date
Classification/Unclassified Controlled Nuclear Information	<input checked="" type="checkbox"/>	<input type="checkbox"/>	D.R. Herman	<i>DR. Herman</i>	4/16/92
Patent - General Counsel	<input checked="" type="checkbox"/>	<input type="checkbox"/>	SILBERGLIN	<i>Silberglin</i>	4/17/92
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PRL Program/Project	<input checked="" type="checkbox"/>	<input type="checkbox"/>		<i>William P. White</i>	4/20/92
Publication Services	<input checked="" type="checkbox"/>	<input type="checkbox"/>	S.A. Forney	<i>Susan A. Forney</i>	4/20/92
Other Program/Project	<input type="checkbox"/>	<input type="checkbox"/>			
Information conforms to all applicable requirements. The above information is certified to be correct.					
References Available to Intended Audience <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No Transmit to DOE-HQ/Office of Scientific and Technical Information <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No Author/Requestor (Printed/Signature) Date D.R. Herman <i>DR. Herman</i> 4/16/92 Intended Audience <input type="checkbox"/> Internal <input type="checkbox"/> Sponsor <input checked="" type="checkbox"/> External Responsible Manager (Printed/Signature) Date <i>E. Yuzis</i> E Yuzis 4/16/92			INFORMATION RELEASE ADMINISTRATION APPROVAL STAMP Stamp is required before release. Release is contingent upon resolution of mandatory comments. <div style="text-align: center;">  </div>		
			Date Cancelled Date Disapproved		

JKE 4/20/92

SUPPORTING DOCUMENT

1. Total Pages 78 77

2. Title Sampling and Analysis Plan of the 284E Area Powerplant Shop Process Wastewater Streams.	3. Number WHC-SD-WM-PLN-034	4. Rev No. Rev 0
5. Key Words SAP, 284E Powerplant, Sampling Plan	6. Author Name: D.R. Herman <i>DR Herman</i> 4/16/92 Signature Organization/Charge Code 55300/55310	
7. Abstract This Sampling and Analysis Plan (SAP) establishes the guidelines for providing data to support a waste designation for each liquid effluent.		
8. PURPOSE AND USE OF DOCUMENT - This document was prepared for use within the U.S. Department of Energy and its contractors. It is to be used only to perform, direct, or integrate work under U.S. Department of Energy contracts. This document is not approved for public release until reviewed. PATENT STATUS - This document copy, since it is transmitted in advance of patent clearance, is made available in confidence solely for use in performance of work under contracts with the U.S. Department of Energy. This document is not to be published nor its contents otherwise disseminated or used for purposes other than specified above before patent approval for such release or use has been secured, upon request, from the Patent Counsel, U.S. Department of Energy Field Office, Richland, WA. <i>Legends removed per request of Wtk Legal</i> DISCLAIMER - This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, nor any of their contractors, subcontractors or their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or any third party's use or the results of such use of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof or its contractors or subcontractors. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.		10. RELEASE STAMP <div data-bbox="1039 1011 1510 1236" style="border: 1px solid black; padding: 5px;"> OFFICIAL RELEASE BY WHC DATE APR 20 1992 STA 4 </div>
9. Impact Level 3		

APPROVED FOR RELEASE
4/20/92

Sampling and Analysis Plan of the 284 E Powerplant Process Wastewater Streams

April 1992



**Westinghouse
Hanford Company Richland, Washington**

**Hanford Operations and Engineering Contractor for the
U.S. Department of Energy under Contract DE-AC06-87RL10930**

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Sampling and Analysis Plan of the 284 E Area Powerplant Process Wastewater Streams

April 1992

Prepared for Westinghouse Hanford Company
Richland, Washington
Under Contract DE-AC06-76RLO 1830

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ABSTRACT

This Sampling and Analysis Plan (SAP) addresses the requirements in the U.S. Department of Energy (DOE) Order 5400.1 (General Environmental Protection Program) and the May 21, 1991, proposed amendments to the Hanford Federal Facility Agreement and Consent Order (Ecology et al. 1989), otherwise known as the Tri-Party Agreement. This SAP provides descriptions of field procedures and methods that will be used to meet the following objectives:

- provide sufficient data to support a waste designation for each liquid effluent
- provide data for potential use in developing Washington Administrative Code (WAC) 173-240 engineering reports for a Washington State Discharge Permit, if needed
- provide sufficient data on chemical and radiological constituents to assist in making an estimate of constituent loading and their potential rate of migration to support an assessment of potential impacts of continued discharge
- support process design of wastewater treatment projects, if necessary.

SUMMARY

The objective of the Sampling and Analysis Plan (SAP) is to address the requirements of the U.S. Department of Energy (DOE) Order 5400.1, General Environmental Protection Plan and the May 21, 1991, proposed amendments to the Hanford Federal Facility Agreement and Consent Order, commonly called the Tri-Party Agreement. Specific requirements addressed in this SAP include

- provide sufficient data to support a waste designation for each liquid effluent
- provide data for potential use in Washington Administrative Code (WAC) 173-240 engineering reports for a Washington State Discharge Permit, if needed
- provide sufficient data on chemical and radiological constituents to assist in making an estimate of constituent loading and their potential rate of migration to support an assessment of potential impacts of continued discharge
- support process design of wastewater treatment projects, if necessary.

This plan describes the 284 E Powerplant. Sample collection is described, including collection methodology, location of collection points, frequency of collection, equipment decontamination, and chain of custody throughout the process. The plan details specific contaminants to be tested and references the specific testing methodology to be used for each. Radiological screening of samples, data management techniques, and quality assurance methods are discussed.

Previous analytical test are summarized in an appendix.

ABBREVIATIONS/ACRONYMS

COD	Chemical oxygen demand
CFR	Code of Federal Regulations
DOE	U.S. Department of Energy
Ecology	Washington State Department of Ecology
EDMC	Environmental Data Management Center
EDTA	Ethylenediaminetetraacetic acid
EMO	Environmental Management Operations
EPA	U.S. Environmental Protection Agency
ESQA	Environmental Services Quality Assurance
gpm	Gallons per minute
HPLC	High-performance liquid chromatography
HPT	Health physics technician
LEMIS	Liquid Effluent Monitoring Information System
MCL	Maximum contaminant level
MCLG	Maximum contaminant level goal
MSDA	Material safety data sheets
NCR	Nonconformance Report
OSM	Office of Sampling and Management
PCB	Polychlorinated biphenyl
RCRA	Resource Conservation and Recovery Act
QA	Quality assurance
QAPP	Quality Assurance Project Plan
QC	Quality control
SAP	Sampling and Analysis Plan
S&ML	Sampling and Mobile Laboratories
SDWS	Secondary Drinking Water Standards
TOC	Total organic carbon
TOX	Total organic halogens
VOA	Volatile organic analysis
WAC	Washington Administrative Code
WHC	Westinghouse Hanford Company

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1.0 INTRODUCTION

This Sampling and Analysis Plan (SAP) addresses the requirements in the U.S. Department of Energy (DOE) Order 5400.1 (General Environmental Protection Program, DOE, 1988) and the May 21, 1991, proposed amendments to the Hanford Federal Facility Agreement and Consent Order (Ecology et al. 1989), otherwise known as the Tri-Party Agreement. It is the policy of DOE and Westinghouse Hanford Company (WHC) to protect the public and the environment through collection of effluent characteristic data that can be compared to the requirements of applicable federal, state, and local regulations.

The SAP provides descriptions of field procedures and methods that will be used to characterize the chemical and radiological constituents of the process wastewater generated by the 284 E Powerplant in the Hanford 200 East Area. This plan includes discussions of the following:

- sampling objectives
- project organization and responsibilities
- site background
- sampling rationale for selecting locations and frequencies
- analytical parameters
- field parameter measurements
- radiological screening
- sample collection methodology
- health and safety
- documentation
- equipment decontamination
- sample custody and transport

- data management
- quality assurance.

The SAP is to be used in conjunction with the Liquid Effluent Sampling Quality Assurance Project Plan (Liquid Effluent QAPP) (WHC 1991b, or the most recent update) and an applicable health and safety plan.

1.1 SAMPLING OBJECTIVES

Overall objectives for the SAP are to

- provide sufficient data to support a waste designation for each liquid effluent
- provide data for potential use in developing WAC 173-240 engineering reports for a Washington State Discharge Permit, if needed
- provide sufficient data on chemical and radiological constituents to assist in making an estimate of constituent loading and their potential rate of migration to support an assessment of potential impacts of continued discharge
- support process design of wastewater treatment projects, if necessary.

The sampling program is based on addressing these objectives and has been tailored based on available information (see Section 2). The results of this sampling program should provide reproducible, representative wastewater characterization. Any amendment to this document will be considered a Class 3 change to the Tri-Party Agreement once this document is approved by the regulatory agencies. Revisions of this document would be under the direction of the facility manager according to standard WHC procedures.

1.2 PROJECT ORGANIZATION AND RESPONSIBILITIES

The program manager will be responsible for coordinating this SAP with the other liquid effluent SAPs prepared under the Tri-Party Agreement and for providing support and technical guidance to the facility manager.

The facility manager will be responsible for overall execution of the project and for environmental compliance. Responsibilities include planning, staffing, scheduling, and coordinating field activities.

The sampling and analysis task leader will be responsible for coordination of sampling and analysis activities, including scheduling operators and health physics technicians (HPT) to support the sampling team, reviewing field logs and sampling techniques, tracking sample chain-of-custodies and data, and seeing that analytical data are filed with the Environmental Data Management Center (EDMC). The task leader will also assist the facility manager with maintaining site-specific sampling schedules, authorizing changes to the sampling and analysis plan, and keeping management informed of potential impacts of schedule deviations on individual and program-wide schedules and budgets.

The sampling team, from Sampling and Mobile Laboratories (S&ML) or a qualified subcontractor, will coordinate directly with operations staff to determine when discharges will be occurring and to schedule sample collection accordingly. Liquid Effluent Monitoring Information System (LEMIS) staff will provide the sample team with samples labels with a unique sample number. WHC will provide sample bottles.

Protocol samples (hereafter called samples) meeting the quality assurance criteria of EPA method or EPA 1986 criteria will be collected by S&ML or the designated sampling team. The sampling team will make a written record of the sampling as discussed in Section 3.3. Once the protocol or baseline samples are taken and analyzed, routine monitoring or process samples specific to the process will be identified.

The Office of Sample Management (OSM) or a qualified contractor will participate in the evaluation and selection of a laboratory or laboratories to perform the needed analysis. These laboratories must meet the criteria of this SAP and the Liquid Effluent QAPP (WHC 1991b).

Environmental samples being sent from the Hanford Site must undergo a radiological screening analysis to monitor radiological activity exceeding the Hanford Site Release Limits. Arrangements will be made with the 222 S laboratory to meet this requirement and to meet the shipping deadline dictated by the shortest analytical holding time listed in Table 2.2 of 24 hours and the shipping schedule at Building 1163. Personnel at Building 1163 will process packaged environmental samples for offsite shipment.

Data resulting from sampling will undergo two levels of review and validation: within the laboratory and outside the laboratory. Initial data reduction, validation, and

reporting at the laboratory will be carried out as described in Section 6.2 of this SAP and the Liquid Effluent QAPP (WHC 1991b). Completed data packages will be validated by OSM, a qualified individual designated by the facility manager, or a qualified subcontractor. The validated data will be stored and managed in a liquid effluent sampling data base. Hard copies of validated data will be retained in the administrative record vault by EDMC.

The U.S. Environmental Protection Agency (EPA) and State of Washington Department of Ecology (Ecology) will have a regulatory oversight role in this project so that the requirements of the Tri-Party Agreement (Ecology et al. 1989) are met. Staff from these regulatory agencies may choose to visit the site, collect split samples during sampling, or visit the analytical laboratory to observe the work being performed.

1.3 SITE BACKGROUND

The Hanford Site is a DOE facility located in Benton County, Washington. The 200 East Area is located in the south-central portion of the Hanford Site, south of the Columbia River. The 284 E Powerplant consists of three facilities: the 282 E Reservoir, the 283 E Water Treatment Plant, and the 284 E Powerhouse (see Figure 1.1). The 284 E Powerhouse is a coal-fired steam plant that provides steam for the 200 East Area operations.

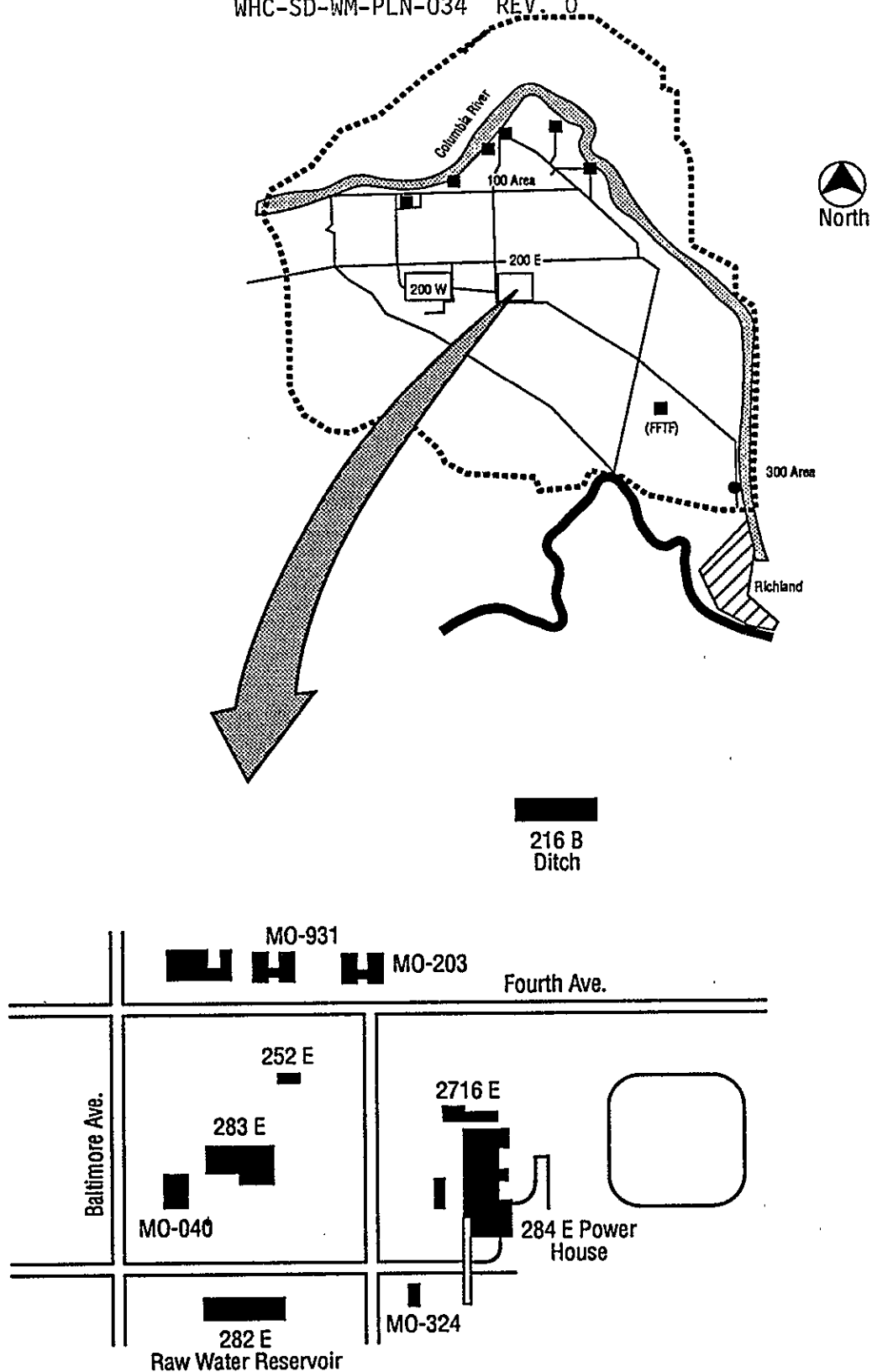
1.3.1 Process Description

Reservoir and Water Treatment Plant

Water drawn from the Columbia River at the 100 B or 100 D Area is pumped to a 25,000,000-gallon reservoir located in the 100 area. Water is pumped from there to the 282 E Reservoir and then pumped to the 283 E Water Treatment Plant (see Figure 1.2). Alum (aluminum sulfate) is added to the water to neutralize electrically charged suspended particles and colloids. The alum-treated water is then sent through a flocculation basin, then into a settling basin. Overflow from the settling basin is filtered through one of four gravity multimedia filters consisting of four layers (ceramic, gravel, sand, and anthracite from bottom to top, respectively). These filters are back-flushed four times a month to remove filtered material or solids. The wastewater flows into the combined wastewater stream that discharges to the 216 B Ditch. The clean, filtered water is then chlorinated and stored in two covered clearwells with a total capacity of 400,000 gallons. This treated water provides potable water to the 200 East Area.

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83-1 RLO3069.MO.SWEHO-Hanford/Aerial View/4-10-92/CS/JH



Source: WHC-EP-0472, 11/7/91, Facility Effluent Monitoring Plan for the 284E and 284W Powerplants, p. 2-3, and personal communications, Ditterman 1992.

Figure 1.1. Aerial View of 284 E Powerplant

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1.6

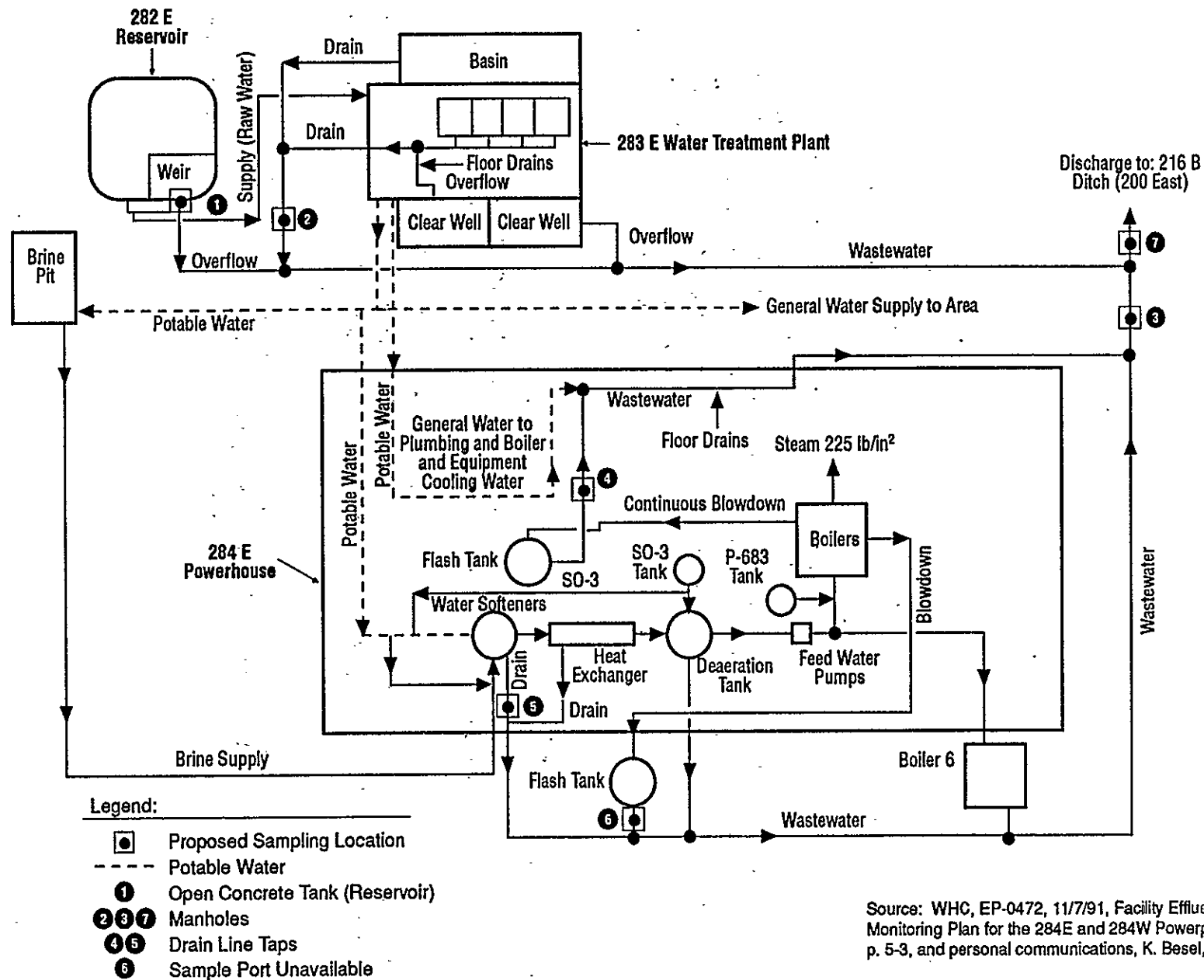


Figure 1.2. Flow Schematic of the 284 E

Powerhouse

Sanitary (potable) water from the clear wells is sent to the 284 E Powerhouse where it is treated with sodium sulfite to destroy residual chlorine and then sent through a water softener to remove mineral hardness, i.e., calcium and magnesium. Polyquest™ 683 is added to the water to control corrosion and scale formation prior to introducing the water into a coal-fired boiler. The generated steam is superheated to between 52°F and 54°F (about 225-pound pressure). The steam generated at the 284 E Powerhouse is used in process facilities in the 200 East Area for heating and process operations (i.e., B Plant, Plutonium-Uranium Extraction Plant, 242 A Evaporator/Tank Farms). The East Powerhouse Complex also provides raw and potable water for process operations, fire protection, and potable water requirements.

A continuous boiler blowdown discharges to the wastewater stream to keep the contaminant concentrations in the boiler water at operational levels. Once per shift, the heavier solids in the bottom of the boiler are discharged to the wastewater stream ("mud drum blowdown"). Approximately twice a week, the water softener is regenerated using a salt brine solution created from water from the potable water supply; the used brine is discharged to the wastewater stream. The salt brine concentration in the zeolite water softener regeneration solution was lowered to 9 percent by weight in late 1989.

1.3.2 Process Changes

Dearborn™ 4846, 4856, and 4812 were previously used at the 284 E Powerhouse to reduce corrosion and scale formation in the boiler. The Dearborn compounds were replaced with Polyquest™ 683 in 1990. Potassium permanganate and barium chloride were used at the 283 W Water Treatment Plant for water treatment and water quality testing in the past. Use of potassium permanganate was discontinued in 1987, and methyl purple indicator replaced barium chloride in 1990. The potential exists for these compounds to appear in the chemical analyses conducted under this SAP, if residuals are present in portions of the piping.

If compounds are detected that are not readily tied to currently used chemicals, comparison of the analytical results to the material safety data sheets (MSDS) for these previously used compounds may aid in the interpretation of the results.

™Dearborn and Polyquest are trademarks of Grace Dearborn.

1.3.3 Wastewater Stream Description

The 284 E Powerhouse, 283 E Water Treatment Plant, and 282 E Reservoir contribute to the common process wastewater discharged to the 216 B Ditch (see Figure 1.2). Table 1.1 lists hazardous chemicals that maybe introduced into the wastewater stream. Table 1.2 presents wastewater contributors, sources, estimated amounts of wastewater contributed by each source, and the approximate frequency of wastewater discharged by each source. Flow rates for wastewater sources at the 284 E Powerhouse will vary seasonally, with higher flow rates due to increased demand for steam heating throughout the 200 East Area during the colder months. Wastewater generated from continuous boiler blowdown, mud drum blowdown, water softener regeneration, and equipment cooling at the 284 E Powerhouse is discharged to the floor trench or directly into floor drains that connect to the 216 B Ditch.

Table 1.1
Hazardous Chemicals Inventory

Product Name	Uses	Potentially Hazardous Compounds
Alum	Flocculent	Aluminum sulfate
Salt	Water softener	Sodium chloride
Chlorine	Disinfectant	Chlorine gas
Lead (in gaskets/valves)	In stored pump gaskets, valve packing	Lead
Mercury (in instruments)	Potentially in stored instruments	Mercury (metallic)
Polyquest™ 683 ^a	Boiler water treatment	<4% Potassium hydroxide
Sulfuric acid	Battery banks	Sulfuric acid
^a Polyquest is a trademark of Grace Dearborn. Source: WHC 1991a, p. 2-12.		

At the facilities listed in Table 1.2, potential sources of dangerous or regulated materials are prevented from entering the effluent stream by using administrative and engineering controls. These controls include lock and tag procedures, source reduction

Table 1.2
Contributors to Flow into 216 B Ditch^a

Contributor	Source	Discharge ^b	Estimated Amount Contributed ^f (flow type)
282 E Reservoir	Cooling water	C	350 gpm ^e
	Pump strainer back flush water	B	
	Heater condensate	C	
	Reservoir overflow	B	7,000 gpm (potable water) ^e
283 E Water Treatment Plant	Filter backwash ^c	B	2,333 gpm./filter x 4 filters ^d x 4 backwash/month (70,000 gal/30 min/ filter x 4 filters x 4 backwash/month)
	Floor drains	B	
	Heater condensate	C	
	Cooling water	C	
	Basin wash down water	B	
	Clearwell overflow	B	
	Basin overflow	B	
	Water testing and sampling station	B	
	Continuous turbidity meter	C	
284 E Powerhouse	Cooling water ^a	C	74 gpm with 2 boilers online (3,250,000 gal/ month)
	Continuous blowdown	C	9 to 21 gpm ^d (varies from 25,000 lb/hr steam load using 540 gal/hr, to 60,000 lb/hr steam load using 1246 gal/hr)
	Mud drum blowdown ^{a,c}	B	2.28 gpm/blowdown (100,000 gal/month)
	Water softener regeneration	B	6.58 gpm (300,000 gal/month)
	Steam heater condensate	C	

^aAll flow estimates taken from WHC 1990.

^bIdentification of discharge type: B = batch, C = continuous.

^cThere are 4 filters in 283 E Water Treatment Plant.

^dNo data available for 284 E Powerplant. Numbers shown are from 284 W Powerplant data. The assumption is made that the powerplants are similar and produce approximately the same flow volumes.

^eBased on information provided by plant operations.

^fFlow quantities not shown are nominal.

(minimization), spill prevention, employee training in hazardous communications, MSDSs, waste handling, and emergency response practices. A detailed description of each effluent contributor follows (WHC 1991).

282 E Reservoir

Water in the reservoir is pumped from the 100 B or 100 D Area River Pump-house. Noncontact cooling water is used to cool the raw water turbine. The cooling water, heater condensate, and reservoir overflow are not expected to contain any dangerous or regulated materials.

282 E Water Treatment Plant

Water pumped from the 282 E Reservoir is treated with alum, filtered, and chlorinated, then stored as a source of potable water in two clearwells. Administrative and engineering controls are used in the plant to prevent sources of water contributing to the effluent stream from coming into contact with dangerous or regulated materials. The resulting effluent stream (listed in Table 1.2) is not considered to be a dangerous waste; however, filter backwash water and basin washdown water are anticipated to be high in suspended solids and metals, including aluminum.

283 E Powerhouse

Sanitary (potable) water from the 283 E Water Treatment Plant is treated with sodium sulfite to destroy residual chlorine and is then passed through a water softener to remove calcium and magnesium prior to introducing the water into the boiler. This aids in minimizing scaling on the tube bundles in the boiler. When the resin in the ion exchange column inside the water softener becomes saturated, the resin is regenerated by passing 9 percent sodium chloride solution through the ion exchange column. Lock and tag procedures are used on control valves so that a concentration of not more than 9 percent sodium chloride is used or discharged to the effluent stream.

Noncontact cooling water is used to cool boiler feed pumps, stoker bearings, water jackets, air compressors, and some fans. Because the cooling water does not come into contact with the equipment, it is not anticipated to be a source of concern.

To minimize potential sources of dangerous or regulated wastes from entering the floor drains located throughout the powerhouse, pump wells (sumps) have been plugged, and also floor drains within 5 feet of any pump have been plugged.

During stream production, minerals not removed in the water softener process collect in the boiler. Two separate blowdown operations are used to remove these minerals: continuous and mud drum. Continuous blowdown is ongoing anytime a boiler is in operation. Mud drum blowdown occurs once per shift to remove solids that have accumulated in the boiler. Blowdown effluent contains antiscaling and oxygen scavenging compounds. Polyquest™ 683 is not considered a dangerous waste after introduction into the process. This is based on the concentration used when this chemical is mixed with water in the mixing tanks prior to entrance into the boiler.

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2.0 SAMPLING LOCATIONS AND FREQUENCIES

2.1 FIELD APPROACH

Sampling locations were chosen based on process knowledge, the configuration of the wastewater discharge piping, and probable accessibility. These sampling locations are also strategically located on the effluent streams from particular operations. The time of sample collection will be directly related to the various operations and a predetermined periodic, sampling schedule. The sampling team will coordinate with operations staff so that the samples are taken at the proper time during the discharges (as described below).

Eight sampling locations have been chosen based on the configuration of discharge lines from the different processes contributing to the overall discharge. A description of these processes and their wastewater streams, a map (Figure 1.1), and a flow schematic of the 284 E Powerplant area showing sampling locations (Figure 1.2) are given in Section 1. The seven sampling locations consist of an open concrete tank (sample location 1, the reservoir), three manholes (sample locations 2, 3, and 7), two drainline sample taps (sample locations 4 and 5), and one drainline which does not currently have a drainline sample tap available (sample location 6). In order to sample mud drum blowdown separately from the other wastewater sources, a drainline sample tap would be needed at sample location 6; given the existing conditions, sample location 3 will be sampled. Sample location 3 is less desirable because it is downstream of sample location 6, after the mud drum blowdown has mixed with other wastewater streams.

The field effort to characterize the 284 E Powerplant wastewater has been phased into four sequential field activities:

- Field Activity 1 will confirm that proposed sample locations are accessible and appropriate for wastewater stream sample collection. In the event that a location does not correspond to the available facility plan, alternative sampling locations will be identified.
- Field Activity 2 tasks will focus on two objectives over two sequential weeks: collecting field measurements and evaluating a combined discharge. During the first week, flow rates and field parameters (conductivity, pH and dissolved oxygen) will be measured for each of the major

wastewater streams and in the common discharge. These data will verify that changes in field parameters (specifically, conductivity) track batch discharges downstream.

Assuming the first week's activities confirm that changes in field parameters result from batch wastewater discharges, then in the second week an attempt will be made to stagger the discharges so that they arrive at sample location 7 at the same time. Field parameters will be used to verify that two sets of wastewater samples were taken during the coincided batch discharges; these samples will be analyzed for a full set of chemical parameters. Chemical results from this chemical sampling, in conjunction with wastestream data, should identify a subset of chemical analytes which are neither expected nor detected in the wastewater process streams. These nondetected analytes will not be quantified in successive sampling.

- Field Activity 3 will extend sampling frequency to weekly events over a one month period. Chemical analyses will be limited to the expected or detected analytes identified in Field Activity 2.
- Field Activity 4 will extend sampling frequency to four sequential quarters. Chemical analyses will be limited the same subset of analytes identified during the initial chemical characterization sampling in Field Activity 2.

Table 2.1 details the specific tasks to be performed during each of the four field activities. The following discussions describe the rationale for the phased approach and for the proposed locations and frequencies.

2.2 FIELD ACTIVITIES

The field effort consists of 4 Field Activities to evaluate the effectiveness of this sampling scheme (see Table 2.1 for sampling details). A schedule for the field effort is shown in Figure 2.1.

Table 2.1
Field Activities and Sampling Frequencies

Page 1 of 4

Field Activity	Data/Samples Collected^a/Frequencies
Field Activity 1	<ul style="list-style-type: none"> • Conduct dye studies to verify the source of wastewater for sample location 2 (see Figure 1.3 and Table 3.1). Release dye into the process piping near the discharge point, upstream from the sample location and see if the dye shows up at the sample location. • Verify access to sample locations 3 and 7.
Field Activity 2, Week 1	<ul style="list-style-type: none"> • Measure flow and field parameters^b only for 7 days. No samples will be collected for chemical analyses. • Collect flow data. • At sample locations 1, 2, 3, and 7 (the reservoir and manholes), use DataSondeTM 3 dataloggers or similar equipment to collect continuous field parameter^b data every 30 minutes. • At sample locations 4 and 5 (drainline sample taps), use portable field instruments to collect manual measurements of field parameters^b every 15 to 30 minutes from 1 hour before batch wastewater discharge begins, continuing through to 1 hour after the batch wastewater discharge ends. If the sample tap is on a continuous discharge (e.g. sample location 4), then samples should be taken every hour. Drainline sample taps should be opened for 1 minute prior to taking the manual measurement (see Section 3.1). • Coordinate with the operations manager to determine when batch wastewater discharges will occur and how long they last. • Examine the field parameter^b data at the end of the week to see if each wastewater batch discharge can be detected by its conductivity and if each batch discharge can be detected at sample location 7 by its conductivity.

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Table 2.1
Field Activities and Sampling Frequencies

Page 2 of 4

Field Activity	Data/Samples Collected*/Frequencies
Field Activity 2, Week 2	<ul style="list-style-type: none"> • Measure flow and field parameters^b for 7 days, as in Field Activity 2, Week 1. Measure these concurrently with the sampling described below. • Determine which wastewater batch discharges can be detected at sample location 7 using conductivity. Those batch wastewater discharges that cannot be detected at sample location 7 will need to be sampled individually (i.e. at sample locations 2, 3, 4, and 5, as needed) for the chemical analysis. • Coordinate directly with operations staff at 282 E Reservoir, 283 E Water Treatment Plant, and 284 E Powerhouse to discharge all continuous and batch wastewater discharges so that the batch discharges reach sample location 7 at the same time. The time lag needed for the wastewater to travel through the piping system can be figured out based on the results of Field Activity 2, Week 1. This combined discharge is the wastewater discharge pulse that will be sampled at sample location 7. • Collect wastewater samples (24-hour composite, if possible, and grab^c) using an ISCO 3700 sampler or similar equipment and a bailer one hour before the pulse or wastewater discharge continuing until one hour after the end of each pulse or wastewater discharge. Analyze samples collected for one hour before and for one hour after the pulse or wastewater discharge for routine parameters only (indicated with an R in Table 2.2). Analyze samples collected during the pulse or wastewater discharge for all the parameters listed in Table 2.2. • Take two rounds of samples during the week. • Review the analytical results to determine which parameters are of interest and the location of the sources of interest.

Table 2.1
Field Activities and Sampling Frequencies

Page 3 of 4

Field Activity	Data/Samples Collected*/Frequencies
Field Activity 3	<ul style="list-style-type: none"> • Measure field parameters at the seven sampling locations during sampling events. • Collect wastewater samples from the sources of interest determined in Field Activity 2, Week 2 (sample location 7 and other sample locations, as discussed above). • Use the appropriate sample collection methodology for the type of location (tank, manhole, drainline sample tap) and sample type (24-hour composite or grab^c). • Collect samples once a week for 1 month (i.e., four rounds of samples) during a single pulse and/or batch wastewater discharge each week (coordinated by the sample team with operations staff as described above). • Samples collected for 1 hour before and for 1 hour after each pulse or batch wastewater discharge will be analyzed for routine parameters (designated with an R in Table 2.2) only. Samples collected during the pulse or batch wastewater discharge will be analyzed for routine parameters and a subset of Appendix IX parameters (chosen on the basis of the analytical results from Field Activity 2, Week 2). • Examine the results to determine which sources of interest need further study. Any parameters not detected during Field Activity 2 and 3 will be eliminated from the parameters list for Field Activity 4.
Field Activity 4	<ul style="list-style-type: none"> • Measure field parameters at the seven sampling locations during sampling events. • Collect one set of wastewater samples once per quarter (for example, January, April, July, and October) for 1 year at the same locations used in Field Activity 3 using the same collection methods.

Table 2.1
Field Activities and Sampling Frequencies

Page 4 of 4

Field Activity	Data/Samples Collected ^a /Frequencies
Field Activity 4 (continued)	<ul style="list-style-type: none"> • Samples collected 1 hour before and 1 hour after a pulse or batch wastewater discharge will be analyzed for any previously detected (in Field Activity 2 and/or 3) routine parameters only. Samples collected during the pulse or batch wastewater discharge will be analyzed for previously detected routine parameters (as described above) and a subset of previously detected (in Field Activity 2 and/or 3) Appendix IX parameters.
<p>^aSample location 6 is the preferred location for sampling mud drum blowdown before it is mixed with other wastewater stream. However, since no sample tap is available at sample location 6, sample location 3 will be used to sample mud drum blowdown after it is combined with other wastewaters from the powerhouse.</p> <p>^bField parameters consist of conductivity, pH, and dissolved oxygen.</p> <p>^cAt manhole locations (1, 2, 3, and 7) grab samples are volatile organics, sulfide, fecal and total coliform, total organic halogens (TOX), and onsite radiation screen samples. At sample tap locations (4 and 5) grab samples are collected for all the parameters listed in Table 2.2.</p>	

2.7

	MONTH															
	1				2	3	4	5	6	7	8	9	10	11	12	13
	WEEK 1	WEEK 2	WEEK 3	WEEK 4												
Field Activity 1 Dye Test	■															
Field Activity 2, Week 1 Field Parameters (7 days, continuous readings)		■														
Field Activity 2, Week 2 Field Parameters and Lab Analysis (2 sets of samples)			■													
Field Activity 3 Field Parameters and Modified Lab Analysis (1 set of samples per week)							■									
Field Activity 4 Field Parameters and Modified Lab Analysis (1 set of samples per quarter)										■		■			■	■

Figure 2.1. Proposed Field Sampling Schedule

2.2.1 Field Activity 1

Because of the age of the facility, some of the available drawings did not adequately indicate whether certain manholes were tied to the discharge lines that need to be sampled. Consequently, some of the chosen sampling locations need to be

checked before samples can be taken. A dye study will be performed to verify that the sample location is a strategic point on the wastewater discharge (e.g. is the manhole chosen for sampling actually located on the discharge line in question?).

Field Activity 1 consists of conducting a dye study in the sewer line to see if the manhole selected as sample location 2 is tied to the discharge line in question. Access to sample locations 3 and 7 will be verified.

2.2.2 Field Activity 2

A review of existing data (see Appendix B) indicated that variations in constituent concentrations could be tracked by measuring the conductivity of the wastewater discharges. In other words, higher chemical constituent concentrations appear to result in higher conductivities. This finding is consistent with knowledge of the wastewater streams that discharge from the 284 E Powerplant area. These wastewater discharges would be expected to have concentrations of salts and solids above background levels. Assuming that the wastewater constituents are fairly consistent from discharge to discharge (a reasonable assumption given the nature of the discharges), the magnitude of the change in conductivity for a given wastewater discharge is expected to be consistent from discharge to discharge over time. Therefore, tracking conductivity should provide a reliable indicator parameter for tracking overall changes in the other constituents of the discharges.

2.2.2.1 Week 1

Because individual wastewater discharges should have different conductivity magnitudes and different batch discharge cycles, the individual batch wastewater discharges should be able to be monitored by measuring conductivity downstream, after several of the discharges come together. Field Activity 2, Week 1 evaluates the effects of the various wastewater discharges on conductivity close to the discharge point (sample locations 2, 4, 5, and 6) and at the "end of pipe" near the 216 B Ditch (sample location 7). Sample location 3 is an intermediate location within the

powerhouse discharge and would probably be used to test mud drum blowdown because proposed sample location 6 does not have a drainline sample tap. Sample location 1 is the background sample location.

In Field Activity 2, Week 1, wastewater flow and field parameters (conductivity, pH, and dissolved oxygen) will be measured for one week during routine operations of the powerplant area. This set of measurements will be used to verify whether conductivity can be used to track the batch discharges. Flow and the field parameters will be measured at, or near, the points of wastewater generation (sample locations 2, 3, 4, 5, and 6). These results will be compared to measurements taken at the "end of pipe" location (sample location 7). If a change in conductivity for each of the wastewater discharges can be detected at sample location 7, then subsequent samples would be justifiably taken from sample location 7 for "end of pipe" samples. Sample locations whose associated discharges cannot be detected and measured at the "end of pipe" would be sampled separately.

During the above evaluation, field parameters will be measured continuously using a data logger (e.g. DataSonde™ 3) at sample locations 1, 2, 3, and 7, where the probe can be continuously submerged. At the other locations, measurements will be taken manually every 15 to 30 minutes because data loggers cannot be used at these locations. The sampling team will coordinate with operations to find out when the batch discharges that must be manually monitored will occur. In the case of the manual monitoring, measurements will be taken one hour before, during, until one hour after batch wastewater discharges take place (see Table 2.1).

2.2.2.1 Week 2

Analysis of the field measurements (flow, conductivity, pH, and dissolved oxygen) should show when the batch discharges were occurring, how long it took a given batch discharge to reach a downstream sampling location, and if the batch discharge can be tracked by conductivity at a downstream sampling location. The field measurements will be evaluated to determine if conductivity can be used as an indicator parameter. The sample locations for chemical analysis in Field Activity 2, Week 2 would be chosen based on how far downstream conductivity can be used to track the individual batch wastewater discharge.

In Field Activity 2, Week 2 of the sampling program, the 284 E Powerplant area processes will be coordinated so that the individual process discharges will arrive at the "end of pipe" sample location 7 at the same time. This combined discharge is

the wastewater discharge "pulse". Field measurements will continue to be taken at sample locations 1, 2, 3, 4, 5, 6, and 7 during Field Activity 2, Week 2 so that conductivity can be used to trace the time the individual wastewaters were discharged.

Samples will be taken before, during, and after the combined discharge. The samples taken during this combined discharge should be representative of the greatest total amount of dissolved solids and highest flow rate that could discharge to the environment at any one time, assuming that operations do not change significantly. During this activity, samples will be taken that are either grab samples or composite samples depending on what is appropriate for a specific sample location or parameter (e.g. all volatile organics samples will be grab samples). The total discharge pulse will be sampled twice at the selected sample locations before, during, and after the discharge so that a representative sample and a "backup" sample are obtained. Analytical data from the two samples will be compared to determine if the samples are consistent. If this sampling does not adequately characterize the wastewaters, more samples will be taken over a longer period of time to more fully characterize the wastewater.

Samples collected before and after each pulse will be sent to an analytical laboratory for routine analyses as indicated with an R in Table 2.2. Routine analysis parameters are those expected to be in the wastewater stream based on available process knowledge and chemical data (see Table 2.2). Samples collected during each pulse will be analyzed for all constituents listed in Table 2.2, including the full 40 CFR 264 Appendix IX list of chemical constituents. A total of six samples (four before and after the pulse for routine analyses and two during the pulse for the entire Table 2.2 list of constituents) would be collected from sample location 7 and analyzed in Field Activity 2, Week 2. Other sample locations may also need to be sampled (see previous discussion). The results will establish a constituent baseline for the combined wastewater discharge. The results also will be used to determine if further testing and evaluation of the individual wastewater sources is needed.

If the analytical results from the combined discharges do not exceed applicable regulatory concentrations, then it is assumed that the combined discharge can be monitored. If the analytical results show concentrations above applicable levels, then additional sampling on those discharge streams showing the highest conductivity may be needed to determine the actual concentration in the wastewater stream causing the exceedences.

Table 2.2
Sample Parameters, Holding Times, Containers, and Preservatives

Page 1 of 2

Parameter	R ^f	Holding Time Extract/ Analysis	Sample Container ^a	Preservative (cool to 4°C)	Method ^b
40 CFR 264 Appendix IX					
Total cyanide		NA/14 days	1/2 gal. plastic	NaOH	EPA 335.2
Sulfide		NA/7 days	500 ml. poly	NaOH + ZnAc	EPA 376.1
Metals ^c	R	NA/6 mos., except Hg 28 days	1 l. poly	HNO ₃ , pH<2	SW 6010/7000 series
Volatile organics (Appendix IX) and ethylene glycol		NA/14 days	40 ml. vial with Teflon™ septum	HCl, pH<2 ^c	SW 8240
Semivolatile organics (Appendix IX) and hydrazine and citrus red No. 2		7/40 days	2 l. amber glass w/Teflon™-lined cap	None ^c	SW 8270
Pesticides/PCBs		7/40 days	2 l. amber glass w/Teflon™-lined cap	None	SW 8080
Organophosphate pesticides		7/40 days	2 l. amber glass w/Teflon™-lined cap	None	SW 8140
Chlorinated herbicides		7/40 days	2 l. amber glass w/Teflon™-lined cap	None	SW 8150
Dioxins/furans		7/40 days	2 l. amber glass w/Teflon™-lined cap	None	SW 8280
Field Parameters					
pH	R	NA/analyze immediately	2 oz. poly or glass	None	SW 9045
Conductivity	R	NA/28 days	250 ml. poly or glass	None	SW 9050
Water Quality					
Total organic carbon (TOC)	R	NA/28 days	250 ml. poly or glass	H ₂ SO ₄ , pH<2	SW 9060
Total organic halogens (TOX)		NA/7 days	500 ml. amber glass w/Teflon™-lined cap	H ₂ SO ₄ , pH<2	SW 9020
Turbidity	R	NA/48 hours	100 ml. poly or glass	None	EPA 180

Table 2.2
Sample Parameters, Holding Times, Containers, and Preservatives

Page 2 of 2

Parameter	R ^f	Holding Time Extract/ Analysis	Sample Container ^a	Preservative (cool to 4°C)	Method ^b
Water Quality					
Fecal coliform		NA/30 hours ^d	100 ml. amber glass	None ^c	SM 908C
Total coliform		NA/30 hours ^d	100 ml. amber glass	None ^c	SM 908A
Alkalinity	R	NA/14 days	250 ml. poly or glass	HCl, pH<2	EPA 310.1
Total phenols		NA/28 days	500 ml. glass	None	EPA 420.1
Chemical oxygen demand (COD)	R	NA/28 days	100 ml. poly or glass	H ₂ SO ₄ , pH<2	EPA 410.1
Fluoride	R	NA/28 days	250 ml. poly	None	EPA 340.2
Chloride	R	NA/28 days	250 ml. poly or glass	None	EPA 325.3
Sulfate	R	NA/28 days	250 ml. poly or glass	None	EPA 375.1
Sulfite	R	NA/24 hours	250 ml. poly or glass	None	EPA 377.1
Nitrate	R	NA/48 hours	250 ml. poly or glass	None	EPA 325.1
Nitrite	R	NA/48 hours	250 ml. poly or glass	None	EPA 354.1
Ammonia	R	NA/28 days	8 oz. poly or glass	H ₂ SO ₄ , pH<2	EPA 350.2
Amenable cyanide		NA/14 days	1 l. poly or glass	NaOH, pH>12 ^c	EPA 335.1
Radiological					
Gross alpha/beta	R	NA/6 mos.	500 ml. poly	HNO ₃ , pH<2	SW 9310
Onsite rad. screen			100 ml. poly	None	
^a Containers should be precleaned glass or polyethylene with necessary preservatives added. Volumes may change based on the requirements of the laboratory performing the analysis. ^b Methods are from EPA 1986, EPA 1983, or APHA 1989. ^c These parameters are sensitive to residual chlorine. If chlorine is present, the volatile and semivolatile organic samples and the fecal and total coliform samples must be preserved with Na ₂ S ₂ O ₃ and the cyanide sample with ascorbic acid. ^d If it is not possible to meet the holding time requirement, the laboratory will be instructed to run these tests as soon as practical after receipt of the sample. ^e Teflon™ is a trademark of E.I. DuPont de Nemours and Company, Inc. ^f R = Routine analysis parameter.					

2.2.3 Field Activity 3

Once conductivity has been established as a characteristic that can be used to track the general quality of the batch discharges and a conductivity baseline has been established for the various discharges, conductivity can be used as the long-term routine monitoring parameter. If conductivity is measured in the field during routine discharges and a conductivity higher than the baseline is seen, it would be easy for the person measuring the discharge to take a sample for laboratory analysis. The constituents for laboratory analysis would be chosen based on the results of the long term sampling in Field Activities 3 and 4.

Field Activity 3 of the sampling plan examines the variability of the field measurements, flow, and detected constituents in the combined wastewater stream. The extent of anomalies, if detected, will be observed by collecting samples and analyzing them for routine analysis parameters (see Table 2.2) as well as a corroborative subset of 40 CFR 264 Appendix IX constituents. The subset of Appendix IX constituents will be chosen based on analytical results from Field Activity 2, Week 2. The samples will be taken before, during, and after process pulses once a week for one month. Samples should be taken on different days of the week during the month to randomize the samples. A total of twelve samples (eight from before and after the pulse for routine analysis parameters and four from during the pulse for a subset of the Table 2.2 constituents) will be collected from sample location 7 and analyzed in Field Activity 3. Other sample locations may also need to be sampled (see Field Activity 2, Week 2). The results then will be evaluated; additional chemical constituents may be removed from the analysis list based on this evaluation.

Field parameters will continue to be taken at sample locations 1, 2, 3, 4, 5, 6, and 7 during Field Activity 3 so that conductivity can be used to trace when each of the individual wastewaters discharged.

2.2.4 Field Activity 4

Field Activity 4 continues the wastewater evaluation by investigating the differences that may occur during the course of the year. Samples will be collected quarterly and analyzed for routine and constituents and the subset of Appendix IX constituents that results from Field Activity 3. Samples should be collected on different days of the week and different weeks in the month to randomize the analytical data. A total of eight samples for routine analysis and four for a subset of the Table 2.2 constituents will be collected from sample location 7 and analyzed in Field

Activity 4. Other sample locations may also need to be sampled (see Field Activity 2, Week 2). Field parameters will continue to be taken at sample locations 1, 2, 3, 4, 5, 6, and 7 during Field Activity 4 so that conductivity can be used to trace when each of the individual wastewaters discharged. As previously stated, this sampling plan is designed to meet the objectives listed in Section 1.1.

9 3 1 2 7 5 3 0 2 5 6

3.0 SAMPLE COLLECTION, IDENTIFICATION, AND HANDLING

Proposed sample locations are listed in Table 3.1.

Table 3.1 Sample Locations ^a	
Location	Sample Point
1	Open concrete reservoir, 282 E Reservoir
2	Manhole, outside 283 E Water Treatment Plant
3	Manhole downstream, 284 E Powerhouse combined effluent
4	Drainline sample tap, 2nd floor 284 E Powerhouse, continuous blowdown
5	Drainline sample tap, ground floor 284 E Powerhouse, water softener blowdown
6	Preferred location for sampling mud drum blowdown; however, sample port unavailable
7	Manhole, above discharge to 216 B ditch
^a Locations shown on Figure 1.2.	

The sampling team will coordinate directly with operations staff to determine when discharge will be occurring and to schedule sample collection accordingly. Doing this will help determine when and if all the processes can be discharged at one time during Week 2 of Field Activity 2.

Wastewater samples and field data will be collected as follows, based on available WHC information (see Table 2.1 for more information):

- Field Activity 1—Perform a dye study to determine if sample location 2 is a valid sampling location, and verify access to manholes at sample locations 3 and 7.
- Field Activity 2, Week 1—Measure field parameters only (pH, conductivity, and dissolved oxygen) at four locations (sample locations 1, 2, 3, and 7) for 1 week. Flow monitoring equipment will be used to record flow data.

- Field Activity 2, Week 2—Samples for chemical analysis will be taken 1 hour before, during, and 1 hour after each combined wastewater discharge pulse. This sampling will occur twice during the week. Samples collected before and after the pulse will be analyzed for routine parameters; samples collected during the pulse will be analyzed for the analytes listed in Table 2.2. Flow data and field parameters will also be collected.
- Field Activity 3—Samples collected for chemical analysis will be taken before, during, and after the combined wastewater discharge pulse. This sampling will happen once a week for 1 month. Samples collected before and after the pulse will be analyzed for routine parameters; samples taken during the pulse will be analyzed for analytes of interest based on the results of Field Activity 2, Week 2. Field parameters will also be collected.
- Field Activity 4—Samples will be collected and analyzed as in Field Activity 3, except that the samples will be collected once per quarter for four quarters to verify the consistency of the discharge.

Field parameters will be monitored continuously using a DataSonde™ 3 or equivalent data logger (assuming there is sufficient water to submerge the data logger probe). The DataSonde™ 3 is a submersible, multiparameter water quality probe and data logger manufactured by Hydrolab Corporation. The probes will be installed in the manholes so that flow is not obstructed. Confined space entry will comply with the Industrial Safety Manual (WHC 1988). Flow data will also be collected and recorded in the field logbook at those sample locations where it can be collected. Sample locations 4 and 5 will be monitored manually for field parameters at specified intervals before, during, and after batch wastewater discharge using portable field monitoring equipment.

3.1 FIELD PARAMETER MEASUREMENTS

The DataSonde™ 3 or similar equipment will be used at four locations (see Table 2.1) to continuously monitor pH, conductivity, and dissolved oxygen. The unit is designed to remain in service for extended periods of time. The DataSonde™ or similar equipment 3 will be field calibrated and checked against standards according to the manufacturer's specifications at the beginning and end of Week 1 of Field Activity 2. Information from the self-contained data logger will be downloaded using a PC communications program (like Procom Plus) to a field computer at the end of the week. Calibration measurements will be recorded in the bound field logbook.

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Samples from the drainline sample taps will be collected 15 to 30 minutes from 1 hour before each batch discharge until 1 hour after each batch discharge. If the sample tap is on a continuous discharge (e.g., sample Location 4), then samples should be taken every hour. Prior to sample collection, the drain line sample taps at 4 and 5 will be opened, and water will be discharged into a container for at least 1 minute to remove any debris or contaminants from the sample taps themselves. A clean cup or beaker will then be filled and used for field parameter measurements. Field parameters will be measured immediately after sample collection. The monitoring probes will be rinsed with a distilled water spray prior to each measurement. The field meters used for these measurements will be calibrated daily in accordance with manufacturers' specifications (which will be kept with each instrument) using appropriate standard solutions. Field parameters and calibration measurements will be recorded in the bound field logbook. Refer to Section 7.0 of the Liquid Effluent QAPP (WHC 1991b) for quality assurance procedures for calibrating field instruments.

3.2 SAMPLE COLLECTION

During Week 2 of Field Activity 2, 24-hour composite samples will be collected at selected sample location(s) using dedicated programmable ISCO™ 3700 samplers or similar equipment. Sample locations will be selected on the basis of field parameter data collected during Week 1 of Field Activity 2. The ISCO™ 3700 sampler(s) can be programmed to collect a specific sample volume at a specific time interval based on process information. ISCO's™ dedicated Teflon™ suction line with an attached stainless steel strainer will be submerged in the wastewater flow at the bottom of the manhole(s) so that flow is not obstructed. The manholes will be monitored for radiation prior to entry. Installing the ISCO™ suction line and strainer in the manhole requires confined space entry and will comply with the Industrial Safety Manual WHC-CM-4-3 (WHC 1988). A 120-volt, 60-Hz power supply is necessary to operate the ISCO™ unit(s).

Grab samples at selected sampling locations will be collected directly from drainline sample taps or with dedicated stainless steel bailers with Teflon™ check valves. Bailers will be lowered on a monofilament line into the wastestream using a cord reel. The full bailer will be retrieved, and a Teflon™ stopcock will be inserted at the check valve so that flow from the bailer can be controlled. Likewise, the drainline sample tap will be used to reduce the flow rate when collecting samples for volatile organic analysis to minimize aeration of the sample and possible volatilization of organic compounds.

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Equipment decontamination procedures and the disposal of contaminated materials is discussed in Section 3.4, Equipment Decontamination.

Field quality control samples will be collected as part of the field sampling effort. Field quality control samples will include equipment blanks, field blanks, trip blanks, and duplicate samples. The analytical schedule for the field quality control samples is shown in Table 3.2. The frequency of quality control sample collection is described below and discussed further in Section 10.0 of the Liquid Effluent QAPP (WHC 1991b).

Table 3.2 Analytical Schedule for Field Quality Control Samples				
Parameter Group ^a	Sample Type			
	Duplicates	Field Blanks	Equipment Blanks	Trip Blanks
Volatile Organics ^a	X	X	X	X
Semivolatile Organics ^a	X		X	
Pesticides and PCBs ^a	X		X	
Herbicides ^a	X		X	
Dioxin/Furans ^a	X		X	
Metals ^a	X		X	
Radionuclides ^b	X		X	
Field Parameters ^b	X			
Miscellaneous Parameters ^c	X		X	
^a Individual constituents in each parameter group are presented in Tables C.1 through C.7 in Appendix C. ^b See Table 2.2. ^c See Table C.7, in Appendix C.				

- **Field Duplicates**—For each phase of sampling activity, a minimum of 5 percent of the total collected samples will be duplicated. Duplicate samples will be retrieved from the same sampling location using the same equipment and sampling technique and will be placed into two sets of identically prepared and preserved containers. All field duplicates will be analyzed independently as an indication of gross errors in sampling techniques.
- **Field Blanks**—Field blanks will consist of pure deionized distilled water, transferred into a sample container at the site, and preserved with the reagent specified for the analytes of interest. Field blanks are used as a check on reagent and

environmental contamination and will be collected at a minimum frequency of 5 percent of the total number of samples taken.

- Equipment Blanks—Equipment blanks will consist of pure high-performance liquid chromatography (HPLC)-grade distilled water washed through decontaminated sampling equipment and placed in containers identical to those used for actual field samples. Equipment blanks are used to verify the adequacy of sampling equipment decontamination procedures and will be collected at a minimum frequency of 5 percent of the total number of samples taken.
- Trip Blanks—Trip blanks consist of pure deionized distilled water added to one or two clean septum-sealed vials, accompanying each batch of containers shipped to the sampling activity. Trip blanks shall be prepared by the laboratory, stored with the samples, and returned unopened to the laboratory. Trip blanks are a check on possible volatile organic compound contamination originating from container preparation methods, shipment, handling, storage, or site conditions. Trip blanks will accompany each shipment of samples scheduled for volatile organic analysis. Trip blanks will be prepared and analyzed at a minimum frequency of 5 percent. However, the project manager or laboratory personnel may elect to analyze additional trip blanks if unusually high or otherwise unexpected concentrations of volatile organics are detected in the field samples.
- MS/MSD—A matrix spike and matrix spike duplicate will be collected from each batch or each 20 samples, whichever is greater, and used for the laboratory's internal quality control/quality assurance checks.

Sample collection during Field Activities 3 and 4 will use similar methodology. See Table 2.1 for details.

3.3 SAMPLE DESIGNATION AND FIELD DOCUMENTATION

Sample bottles will be tracked from the point of sample origin to the laboratory in accordance with a chain-of-custody system described in Section 3.5, Sample Custody and Transport. LEMIS will provide prepreserved sample bottles labeled with a unique sample number. The general numbering method will be as follows:

XX - nnnn

where

XX = Sample point designation

nnnn = Computer-generated sequential serial number

The bottles will be labeled with these numbers. Also, each bottle will be identified with a bar code sticker attached to the bottle by the bottle manufacturer. The bar code will identify the bottle lot number and individual bottle number. Additional information recorded on the label will include:

- time sample was collected to the nearest minute using a 24-hour clock (military time system)
- analysis
- preservative
- customer
- sampler's initials and name printed
- type of sample (i.e., grab or composite).

The sampling team shall maintain a written record of sampling activities and field observations in a bound field logbook. All logbook entries shall be completed in non-erasable ink. Any required corrections to the information in the logbook will be made by drawing a line through the erroneous information, entering the correct information, and initialing and dating the change. The erroneous information should remain legible.

At a minimum, the following information should be noted in the bound field logbook:

- all information required under Section 6.1 of the Liquid Effluent QAPP (WHC 1991b)
- date, time, and sample identification number for each sample collected
- type of sample being collected (i.e., grab or composite)
- scheduled analyses for each sample collected
- qualitative indication of sample turbidity and color
- field parameter measurements and time they were measured

- lot numbers and expiration dates for calibration solutions and gases
- equipment manufacturer, model number, and serial number
- names of sampling personnel and HPT
- radiological screening results for each sample
- deviations from procedure described in this SAP
- daily signatures for each person making logbook entries
- sketches, drawings
- any other pertinent information.

During field activities the field logbook will be kept under the control of the field team. Upon completion of the field effort, the field logbook will be managed in accordance with QR 17.0, Quality Assurance Records (WHC 1989).

3.4 EQUIPMENT DECONTAMINATION

All metal and glassware used in the sample collection that are not certified pre-cleaned will be decontaminated prior to first use, then dedicated to a specific sampling point [according to Resource Conservation and Recovery Act (RCRA) Groundwater Technical Enforcement Guidance Document, 1986 (referenced in Chapter 11 of EPA 1986)]. Decontamination will consist of

- trisodium phosphate and tap water wash
- deionized water rinse
- isopropyl alcohol rinse
- deionized water rinse
- nitric acid rinse (10 percent solution)
- HPLC-grade water.

The isopropyl alcohol, nitric acid, wash water, and deionized water rinsate will be collected in separate labeled containers for disposal according to applicable regulations.

As discussed previously in Subsection 3.2, Sample Collection, an equipment blank will be collected by conducting the third rinse with HPLC-grade organic-free water and collecting the rinsate for analysis.

Field meter sensors (e.g., pH, dissolved oxygen (D.O), and specific conductance probes) used to monitor the process wastewater will be rinsed with deionized water after measurement.

At the end of each sampling event, all equipment exposed to the process wastewater will be decontaminated by washing in a mild trisodium phosphate solution and rinsing with deionized water. This washwater will be disposed of according to applicable regulations.

3.5 SAMPLE CUSTODY AND TRANSPORT

Samples will be routed to the selected contractor or subcontractor laboratory for analysis consistent with Section 8.0 of the Liquid Effluent QAPP (WHC 1991b). Samples of the 200 Area process wastewater will be shipped by overnight courier to a designated analytical laboratory.

All samples will be packaged for shipment in iced coolers. Radiological screening of a representative portion of each sample delivery group will be conducted by the laboratory in Building 222 S (see Section 5 for more information on radiological screening). A chain-of-custody record will be completed to accompany each cooler shipped or delivered to the laboratory from the field. (A copy of a typical chain-of-custody record is provided in Appendix A.) At a minimum, the following information will be provided on the chain-of-custody record by the sampling team:

- all information required under Section 6.1 of the Liquid Effluent QAPP (WHC 1991b)
- chain-of-custody number
- project name and number
- client name
- project manager

- sampler's name and title
- sample locations
- sample identification
- date and time of sample collection
- type of sample (i.e., grab or composite)
- requested analyses
- number of containers
- type of container, preservative, and sample volume
- signatures of all persons having custody of the sample from collection until receipt by the laboratory
- inclusive dates and times of sample possession
- courier name and airbill number (remarks column).

In addition, the laboratory identification number(s) will be entered on the chain-of-custody record by the laboratory sample custodian when the samples arrive at the laboratory. A copy of the signed chain-of-custody record with laboratory assigned identification numbers written in the appropriate column should be returned to OSM.

The chain-of-custody record will be completed in ink. Any required corrections to the information provided on the chain-of-custody record will be made by drawing a line through the erroneous information, entering the correct information, and initialing and dating the change. The erroneous information should remain legible. Any unused sections of the form will have zig-zag lines drawn through them to indicate information is not missing.

The original signature copy of the chain-of-custody record will be enclosed in a self-sealing plastic bag with the Offsite Property Control Form and Total Activity Report (see Section 5) and secured to the inside of the cooler lid. Adhesive labels on the sample bottles will be completed in waterproof ink and covered with clear plastic tape. The glass sample containers will be packaged in bubble wrap or in customized foam packing to protect them from accidental breakage during shipment. Sample bottles will be placed in individual self-sealing bags to contain any potential leakage and then placed in

a cooler. Ice (packaged in self-sealing bags) will be placed around the sample containers to maintain samples at approximately 4°C during shipment to the laboratory. Remaining cooler space will be filled with bubble wrap. A copy of the chain-of-custody record will be retained by the sample team leader and placed in the bound field logbook. The HPT in the shipping and receiving area of Building 1163 will monitor each cooler for alpha, beta, and radon radiation of the external packaging prior to release to a courier. Final sample package preparation will include:

- sealing the drain plug and lid seam with waterproof tape
- attaching a minimum of two chain-of-custody seals in a way that they would be broken if the cooler was opened
- attaching a shipping label
- attaching the WHC Offsite Property Control Form and the courier's shipping papers to the lid.

As an identifying measure, each cooler will be given a unique name written with waterproof ink on the top and side of the cooler. Each time a cooler changes possession, both the person relinquishing and the person accepting custody must sign and date the chain-of-custody record. As long as the custody record is sealed inside the sample cooler and chain-of-custody seals remain intact, representatives of courier companies will not be required to sign the custody record. Shipping papers provide documentation of custody for the courier company.

A sample team member will telephone the laboratory each time a sample delivery group is shipped. The laboratory will be informed of the number of samples that will be arriving, the expected arrival time, and the analyses that will be required. Laboratory notification will be documented in the bound field logbook by the team member making the telephone call. The sample custodian receiving the samples at the laboratory shall sign and date the chain-of-custody record to acknowledge receipt of the samples. Once the samples are received at the laboratory, laboratory personnel will be responsible for maintaining internal logbooks and records that document sample custody throughout sample preparation and analysis. Further details on laboratory chain-of-custody are provided in the Liquid Effluent QAPP (WHC 1991b). The sample team leader will transmit all shipping documentation (i.e., chain-of-custody record, Total Activity Report and Off-site Property Control Form, courier, and airbill number) by facsimile to OSM or a designee, within 24 hours of sample shipment. The laboratory will transmit by facsimile to OSM or a designee copies of the shipping documentation with the laboratory receipt signature within 24 hours of receiving the samples. OSM or the designee will inform the sample team leader that it was received.

4.0 ANALYTICAL PARAMETERS AND PROCEDURES

The chemical parameters for analysis include the groundwater monitoring list (40 CFR 264, Appendix IX), three Hanford specific parameters (hydrazine, ethylene glycol, and citrus red No. 2) and water quality parameters as required by the Liquid Effluent QAPP (WHC 1991b). Table 2.2 lists the proposed analytical parameters. Tables C.1 through C.6 in Appendix C list specific Appendix IX parameters and the target detection limits (based on EPA 1986) for these parameters. Table C.7 lists target detection limits for the other parameters. The target detection limits may be redefined after final laboratory selection by OSM or approved designee. Actual detection limits depend on the nature of the matrix and will be reported for each parameter. The number of analytes may be reduced after a baseline is established (e.g., after the results Week 2 of Field Activity 2, of or Field Activity 3 are analyzed).

The analytical methodology will be based on standard EPA methods, the methodology has been described in the Liquid Effluent QAPP (WHC 1991b). Standard method references for the parameters are listed in Table 2.2. The analytical quality control procedures will follow the individual EPA analytical methodology and the Liquid Effluent QAPP (WHC 1991b) (see Section 10.0).

5.0 RADIOLOGICAL SCREENING

Environmental samples being sent from the Hanford site must undergo a radiological screening analysis to monitor for radiological activity exceeding the Hanford Site Release Limits. This requirement applies to all samples collected as part of this project. Arrangements will be made with the 222 S laboratory to meet this requirement and to meet the shipping deadline dictated by the shortest analytical holding time of 24 hours (see Table 2.2) and the shipping schedule at Building 1163. Note that the sulfite holding time is 24 hours; fecal and total coliform holding times are 30 hours.

A HPT will field-screen the samples prior to their leaving the sample area. In addition, approximately 100 ml of the wastewater sample from each sample point will be collected in plastic sample containers and delivered to the laboratory at Building 222 S. The analytical results (the Total Activity Report) will be transmitted from the laboratory by facsimile machine to a predetermined Hanford site building where the field sample team will pick it up. If the sample exhibits radioactivity at levels below the Hanford Site Release Limits, the sample team may take the samples (packaged in a cooler and ready for shipment) in accordance with applicable Department of Transportation regulations (49 CFR) to the Hanford Shipping and Receiving office at Building 1163 in the 1100 Area. In order for the samples to leave the site on the same day as collected, the samples and the radiological screening results must arrive at Building 1163 by 1:30 p.m. to allow adequate time for shipping and receiving personnel to prepare and sign an Offsite Property Control Form. The resident HPT in the shipping and receiving area will monitor the sample packaging for radiation prior to its release to an overnight courier. Copies of the Total Activity Report and Offsite Property Control Form must accompany the chain-of-custody record in the cooler (see Section 3.5). Copies of all three will be retained by the sample team leader and placed in the field logbook and sample file.

6.0 DATA MANAGEMENT AND QUALITY ASSURANCE

6.1 PACKAGE DATA

The laboratory data package will include the deliverables described in EPA (1986) Section 1.5, including all appropriate quality control (QC) summary sheets. The laboratory will not submit the chromatograms (Section 1.5.9) or the mass spectra (Section 1.5.11), but they will retain these documents as described in the Liquid Effluent QAPP (WHC 1991b). The full documentation will be transferred to WHC at the end of the project as described in the Liquid Effluent QAPP (WHC 1991b) Section 9.1. Validated data for nonradioactive samples will be available to regulators within 86 days after sampling as indicated in Section 15 of the Liquid Effluent QAPP (WHC 1991b). Validated data for low-level samples will be available to the regulators no later than 126 days after the start of sampling.

6.2 DATA REDUCTION, VALIDATION, AND REPORTING

Initial data reduction, validation, and reporting will be carried out by the laboratory per specific standard method guidelines. The laboratory will provide documentation per quality assurance project plan (WHC 1991b) requirements, to include quality control data.

Data will also be reviewed outside the laboratory by OSM personnel, a qualified individual designated by the facility manager, or qualified contractor independent of the laboratory. The data review validation process is described in the Liquid Effluent QAPP (WHC 1991b).

6.3 PERFORMANCE AND SYSTEM AUDITS

Environmental Services Quality Assurance (ESQA) personnel will monitor adherence to the quality control procedures and conduct audits as deemed necessary to fulfill the requirements specified in the Liquid Effluent QAPP (WHC 1991b). This may include field and laboratory visits to evaluate the execution of sampling, sample control, chain-of-custody procedures, recordkeeping, sample handling, and laboratory quality control procedures. The cognizant QA manager will monitor work product quality to evaluate whether the project is performed in accordance with approved procedures and instructions.

QA, EPA, and Ecology personnel may also elect to visit the sampling site and laboratories to audit conformance with the procedures specified in this document or the Liquid Effluent QAPP (WHC 1991b).

6.4 PREVENTIVE MAINTENANCE

Each laboratory is responsible for the maintenance of its analytical equipment. Preventive maintenance for field equipment will be carried out in accordance with the manufacturer's recommended procedures and schedules. The manufacturer's operating and calibration guidelines for field instruments are available in the operations and maintenance manuals for the specific equipment. Spare parts can include batteries, replacement check valve and ball for bailers, and spare pH, dissolved oxygen, and conductivity probes. A suitable inventory of spare parts shall be maintained, as recommended by the manufacturer.

6.5 CORRECTIVE ACTION

If quality control audits result in the detection of unacceptable conditions or data, the facility manager or sample task leader will be responsible for developing and initiating corrective action. The cognizant QA manager will be notified if nonconformance will compromise the intended use of the data. Corrective action may include:

- reanalyzing samples if holding-time criteria permit
- resampling and analyzing
- evaluating and amending sampling and analytical procedures
- accepting data with an acknowledgment of the level of uncertainty.

Corrective action for field measurements may include:

- repeating the measurements
- checking instrument adjustments to see that they are appropriate for the ambient conditions (such as temperature)
- checking the batteries

- checking the calibration
- replacing the instruments or measuring devices.

Corrective action and any deviations from the procedures specified in this SAP or the Liquid Effluent QAPP (WHC 1991b) will be documented on the appropriate Non-conformance Report (NCR) form and submitted to the facility manager for approval, as required by the Liquid Effluent QAPP (WHC 1991b).

9 3 1 2 7 5 3 0 2 7 1

7.0 REFERENCES

40 CFR 264 Appendix IX
49 CFR

APHA-AWWA-WPCF. 1989. Standard Methods for the Examination of Water and Wastewater, 17th edition. Washington, D.C.

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Westinghouse Hanford Company (WHC). 1991b. Liquid Effluent Sampling Quality Assurance Project Plan. WHC-SD-WM-QAPP-011. Rev 0. September. Richland, Washington.

APPENDIX A

TYPICAL CHAIN-OF-CUSTODY FORM

9 3 1 2 7 3 3 0 2 7 3

APPENDIX B

EVALUATION OF AVAILABLE DATA

9 3 1 2 7 5 3 0 2 7 5

APPENDIX BEVALUATION OF AVAILABLE DATA

To develop a defensible sampling strategy and rationale, the available existing data had to be reviewed and evaluated. WHC sampled wastewater from the 284 E Powerplant, including water softener regeneration and mud drum blowdown and specific streams constituting batch input to the continuous powerplant discharge. Analytical data of detected analytes, tabulated in the Operations Support Services Department August 1990 report, "284 E Powerplant Wastewater Stream-Specific Report" (WHC 1990), have been evaluated in preparation of the proposed sampling program. Table B.1 summarizes the sampling dates and count of detected analytes in each of the three streams across the chemical classes of inorganic/metals, radioactivity, organic compounds, including aggregated measurements and individual quantification of volatiles, and water quality.

Samples from the two batch blowdown streams and from the continuous blowdown were collected, independently, over three partially overlapping time periods. Mud drum blowdown was sampled between September 1985 and August 1989. Parameters quantified and detected represented between 1 and 6 sampling events over this time frame and were dominated by inorganics and metals and the two measures of gross radioactivity. Limited subsets of organic (aggregate and volatile analytes) and water quality (4 analytes) parameters were detected over the period the mud drum blowdown was sampled. Semivolatiles were either not quantified or not detected in the mud drum blowdown discharge stream. Beginning in November 1987, a limited subset of parameters were quantified and detected from samples collected from the water softener regeneration stream. Between November 1987 and May 1988, analytes were quantified from a single to a maximum of eight samples. Sampling of the continuous blowdown was begun in August 1988 and continued through March 1990. The set of analytes that were quantified and detected was more extensive, particularly in terms of water quality parameters. Sampling and/or detection frequencies from continuous blowdown samples varied among parameters. The majority of detected parameters were represented by eight or more observations over the 19-month period.

Inspection of available results from the continuous blowdown indicated no obvious increasing or decreasing trends in parameter concentrations through the period of monitoring. In general, chemical concentrations and radiological activity measured in the mud drum blowdown discharge stream were comparable to those reported from the continuous blowdown discharge. While only a single sample was collected during the period of overlap of the two streams sampling events, both streams exhibited consistent

Table B.1
Westinghouse 284 E Powerplant Sampling
Data Summary

Page 2 of 2

Sample Dates	08/88 - 03/90	11/87 - 05/88	09/85 - 08/89
Analytical Class	Number of	Number of	Number of
Analytical Parameter	Samples	Samples	Samples
	Continuous	Softener	Mud Drum
	Blowdown	Regeneration	Blowdown
Vanadium	8		6
Alkalinity	4		
Chromium	4		
Phosphate			7
Sulfide			6
Radioactivity			
Alpha Activity	7	1	4
Beta Activity	7	4	6
Organic			
TOX (as Cl)	8	4	6
TOC	8		6
Total Carbon	4		
Volatile Organic (VOA)			
Acetone	15	1	5
Trichloromethane	19	8	12
1-Butanol	7		2
Dichloromethane	19		12
2-Butanone	19		
Tetrahydrofuran	11		
1,1,1-Trichloroethane		8	
Chloromethane			12
Water Quality			
Conductivity	8	4	6
Total Dissolved Solids (TDS)	4	4	1
pH	8	4	
Suspended Solids	4		
Temperature	8		6
Ignitability	4		1

Table B.1
Westinghouse 284 E Powerplant Sampling
Data Summary

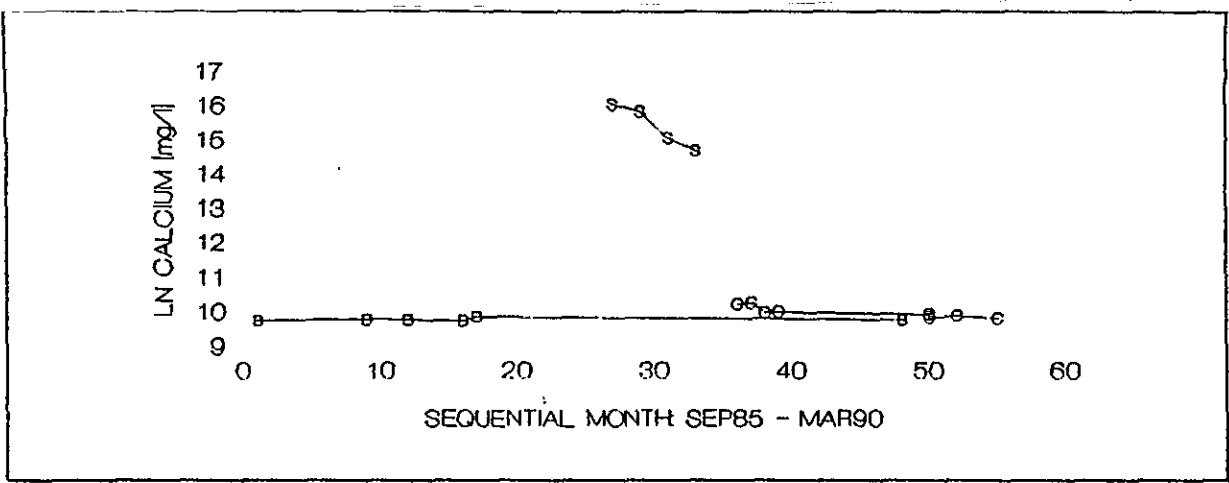
Page 1 of 2

Sample Dates	08/88 - 03/90	11/87 - 05/88	09/85 - 08/89
Analytical Class Analytical Parameter	Number of Samples Continuous Blowdown	Number of Samples Softener Regeneration	Number of Samples Mud Drum Blowdown
Inorganic/Metal			
Aluminum	9	4	5
Ammonia	9	4	5
Barium	13	4	5
Cadmium	13	4	5
Fluoride	18	8	
Iron	8	4	5
Lead	13	4	3
Magnesium	9	4	5
Manganese	9	4	5
Potassium	9	4	5
Sodium	9	4	5
Strontium	9	4	
Sulfate	9	4	5
Uranium	8	4	5
Zinc	8	4	6
Calcium	8	4	
Chloride	8	4	
Copper	8	4	
Arsenic	4		1
Boron	4		1
Mercury	4		7
Nitrate	8		6
Reactivity Cyanide	4		1
Reactivity Sulfide	4		1
Selenium	4		1
Silicon	4		1
Silver	4		1

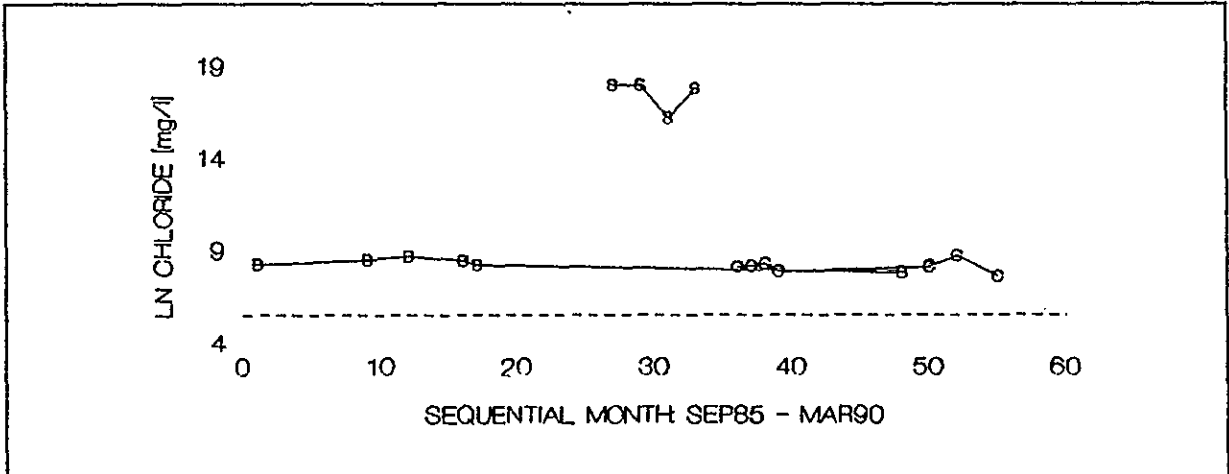
concentration levels (in most contaminants) before (mud drum) and following (continuous) the period of sample overlap. Contaminant levels in the water softener regenerate discharge stream were markedly higher than levels in either the mud drum blowdown or the continuous blowdown discharge. The extent to which these elevated levels of water softener regenerate impact the total discharge cannot be evaluated until simultaneous samples are collected in the two streams during water softener regenerate blowdown release.

Figures B.1 and B.2 present time series of a subset of parameters: calcium, chloride and conductivity, and gross alpha radiation, gross beta radiation and conductivity, respectively. The time axis represents the time period between the first mud drum blowdown sample collected (September 1985) and the final continuous blowdown discharge sample collected (March 1990), a period of approximately 60 months. The axis is divided into sequential months. The points have been coded B for mud drum blowdown, S for water softener regeneration and C for continuous blowdown discharge sample results. Data have been transformed to natural logs to facilitate interpretation. Because the ditch may be recharging groundwater, the Washington groundwater standards (WAC 173-200) were compared to existing data. WAC 173-200-040(2)(b) states that the most stringent criteria should be used for: groundwater quality criteria, federal maximum contaminant level goals (MCLGs), federal maximum contaminant levels (MCLs), or the state MCLs. Criterion for chloride is the Washington State Secondary Drinking Water Standard (SDWS) MCL. The gross alpha and gross beta criteria are the Washington State Groundwater Quality Criteria. Where Washington criteria exist, the criterion level has been indicated with a broken line. The two figures illustrate the trends described above and indicate levels that: 1) consistently exceed Washington criteria in all streams sampled (e.g., chloride), 2) occasionally exceed criteria in some streams (e.g., gross beta activity), 3) are consistently below criteria in all streams sampled (e.g., gross alpha activity).

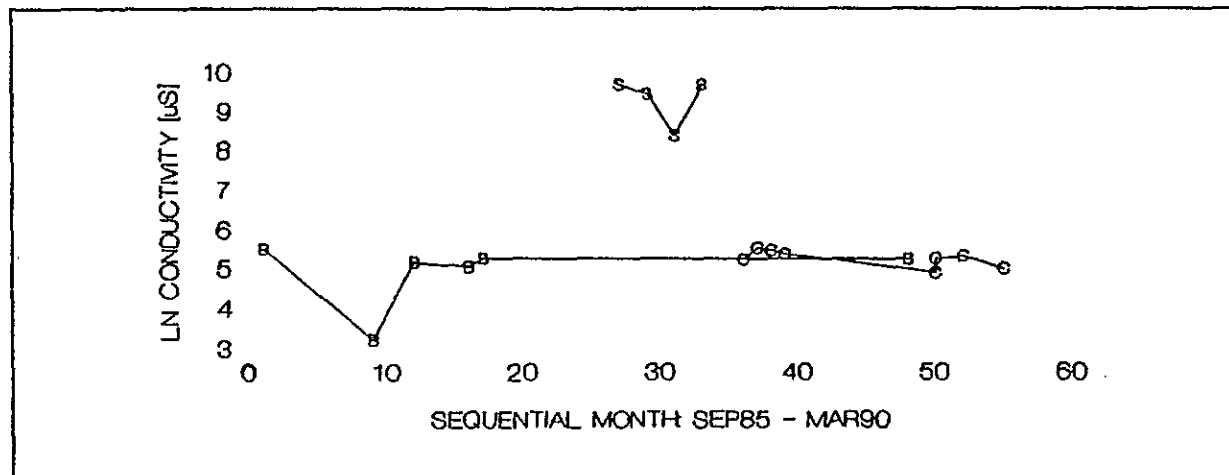
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284-E DISCHARGE CALCIUM [LN] NO CRITERION



284-E DISCHARGE CHLORIDE [LN] ---- WASHINGTON 2° MCL



284-E DISCHARGE CONDUCTIVITY [LN] NO CRITERION

FIGURE B.1, 284 E Powerplant Monitoring Data: September 1985 - March 1990, Conductivity/Water Quality

- LEGEND**
- B Mud drum blowdown
 - S Water softener regeneration
 - C Continuous blowdown
 - pCl/l picoCurles per liter
 - µS microSiemens

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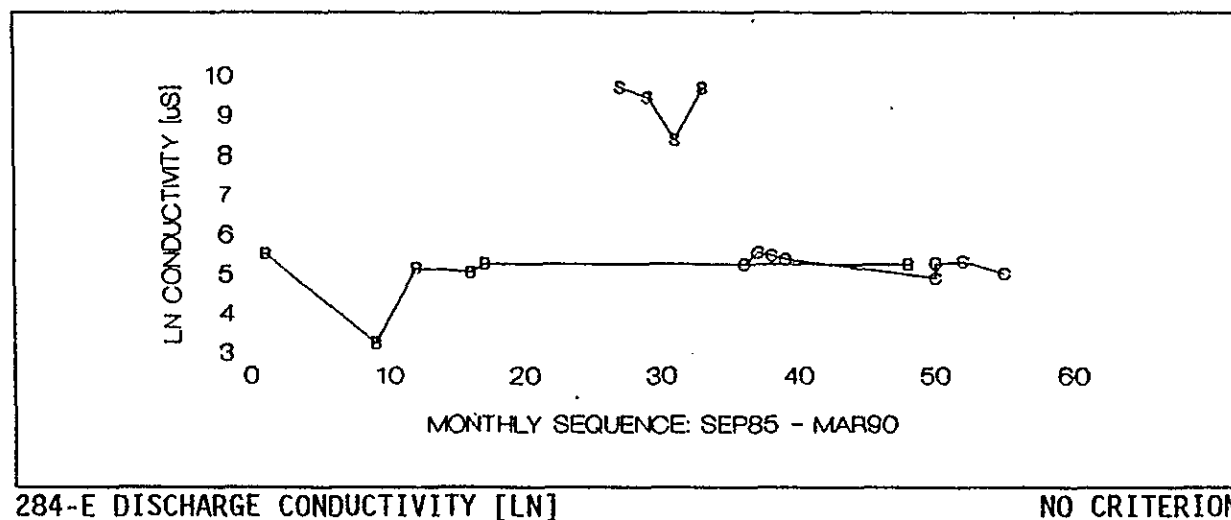
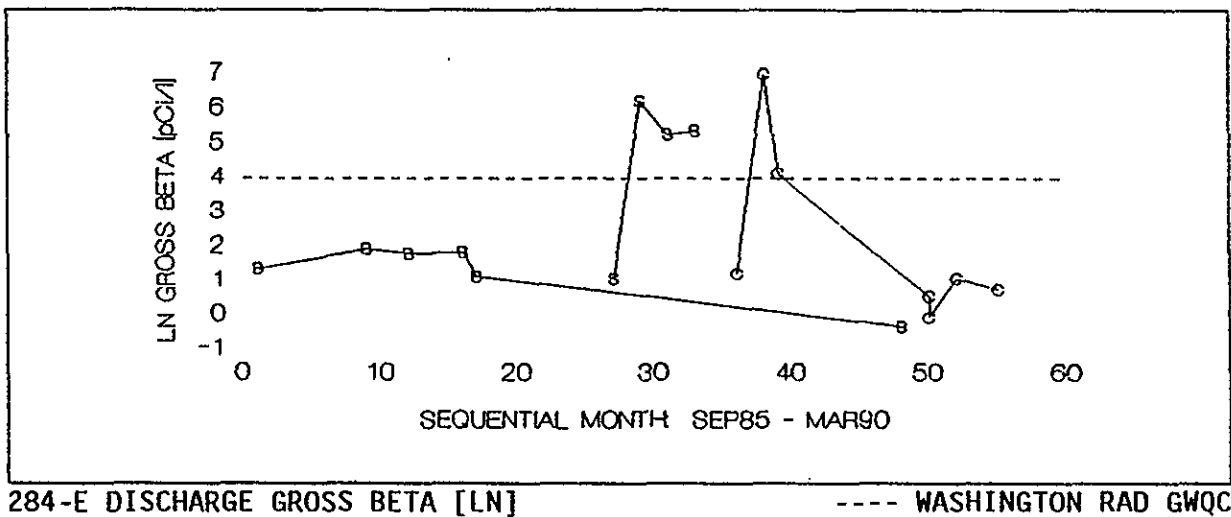
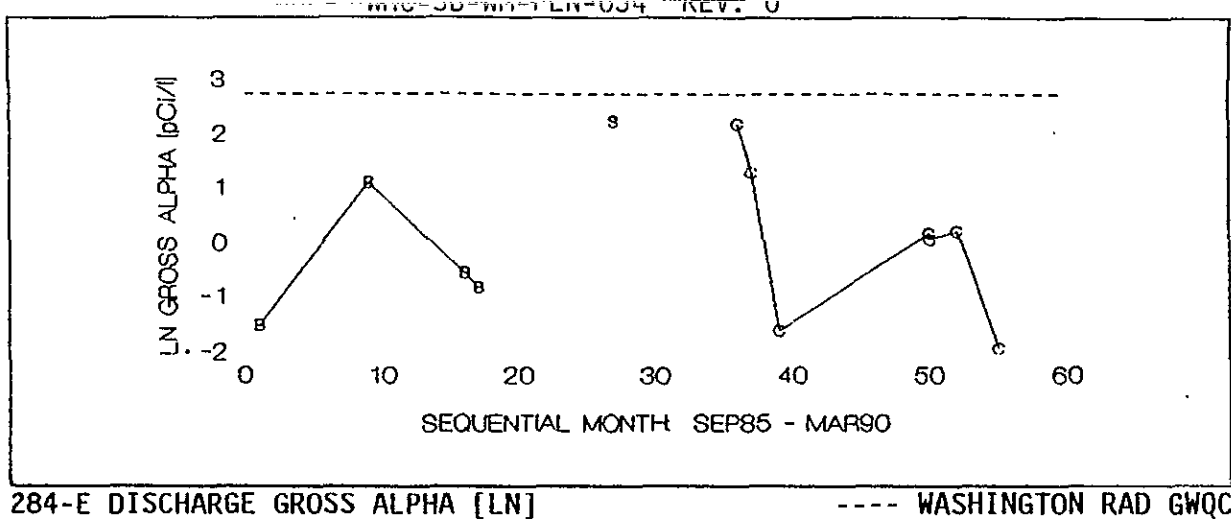


FIGURE B.2. 284 E Powerplant Monitoring Data: September 1985 – March 1990. Conductivity/Radiological Activity

LEGEND

- B Mud drum blowdown
- S Water softener regeneration
- C Continuous blowdown
- pCi/l picoCuries per liter
- μS microSiemens

APPENDIX C

ANALYTICAL PARAMETERS

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Table C.1
40 CFR 264 Appendix IX Volatile Organics
Method 8240^a

Page 1 of 3

Parameter	Target Water Quantitation Limit (µg/l)
Acetone	25
Acetonitrile	100
Acrolein	100
Acrylonitrile	100
Allyl chloride	5.0
Benzene	5.0
Bromodichloromethane	5.0
Bromoform	5.0
Bromomethane (methylbromide)	10
Carbon disulfide	5.0
Carbon tetrachloride	5.0
Chlorobenzene	5.0
Chloroprene	5.0
Chloroethane	10
Chlorodibromomethane	5.0
Chloroform	5.0
Chloromethane	10
DBCP (1,2-dibromo-3-chloropropane)	5.0
EDB (1,2-dibromoethane)	5.0
Dibromomethane	5.0

Table C.1
40 CFR 264 Appendix IX Volatile Organics
Method 8240^a

Page 2 of 3

Parameter	Target Water Quantitation Limit (µg/l)
trans-1,4-Dichloro-2-butene	5.0
Dichlorodifluoromethane	10
1,1-Dichloroethane	5.0
1,2-Dichloroethane	5.0
trans-1,2-Dichloroethene	5.0
1,1-Dichloroethene	5.0
Dichloromethane	25
1,2-Dichloropropane	5.0
cis-1,3-Dichloropropene	5.0
trans-1,3-Dichloropropene	5.0
1,4-Dioxane	100
Ethyl benzene	5.0
Ethyl cyanide	5.0
2-Hexanone	10
Iodomethane	5.0
Isobutyl alcohol	100
Methacrylonitrile	5.0
Methyl ethyl ketone (2-butanone)	25
4-Methyl-2-pentanone	10
Styrene	5.0

Table C.1
40 CFR 264 Appendix IX Volatile Organics
Method 8240^a

Page 3 of 3

Parameter	Target Water Quantitation Limit (µg/l)
1,1,1,2-Tetrachloroethane	5.0
1,1,2,2-Tetrachloroethane	5.0
Tetrachloroethene	5.0
Toluene	5.0
1,1,1-Trichloroethane	5.0
1,1,2-Trichloroethane	5.0
Trichloroethene	5.0
Trichlorofluoromethane	5.0
1,2,3-Trichloropropane	5.0
Vinyl acetate	10
Vinyl chloride	10
Total xylenes	5.0
^a Method 8240 must be modified to include all the listed parameters. The method is from EPA 1986.	

Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270^a

Page 1 of 6

Parameter	Target Water Quantitation Limit (µg/l)
Acenaphthene	10
Acenaphthylene	10
Acetophenone	10
2-Acetylaminofluorene	10
4-Aminobiphenyl	10
Aniline	10
Aramite	10
Benzo(a)anthracene	10
Benzo(b)fluoranthene	10
Benzo(k)fluoranthene	10
Benzo(g,h,i)perylene	10
Benzo(a)pyrene	10
Benzyl alcohol	20
4-Bromophenyl phenyl ether	10
Butyl benzyl phthalate	10
2-sec-Butyl-4,6-dinitrophenol (Dinoseb)	10
4-Chloroaniline	20
4-Chloro-3-methylphenol	20
2-Chloronaphthalene	10
2-Chlorophenol	10
4-Chlorophenyl phenyl ether	10

Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270^a

Page 2 of 6

Parameter	Target Water Quantitation Limit (µg/l)
o-Cresol	10
m & p-Cresol(s)	10
Chrysene	10
Dibenzo(a,h)anthracene	10
Dibenzofuran	10
Di-n-butyl phthalate	10
1,2-Dichlorobenzene	10
1,3-Dichlorobenzene	10
1,4-Dichlorobenzene	10
3,3'-Dichlorobenzidine	20
2,4-Dichlorophenol	10
2,6-Dichlorophenol	10
Diethyl phthalate	10
Dimethoate	--
p-Dimethylaminoazobenzene	10
7,12-Dimethylbenzanthracene	10
3,3'-Dimethylbenzidine	10
a,a-Dimethylphenethylamine	10
2,4-Dimethylphenol	10
Dimethyl phthalate	10
1,3-Dinitrobenzene	10

Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270"

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Parameter	Target Water Quantitation Limit (µg/l)
4,6-Dinitro-o-cresol	50
2,4-Dinitrophenol	50
2,4-Dinitrotoluene	10
2,6-Dinitrotoluene	10
Di-n-octyl phthalate	10
Diphenylamine	10
Disulfoton	--
Dimethoate	--
bis(2-chloroethyl)Ether	10
bis(2-chloroisopropyl)Ether	10
Ethyl methacrylate	10
Ethyl methanesulfonate	10
Ethyl parathion	--
Famphur	--
Fluoranthene	10
Fluorene	10
Hexachlorobenzene	10
Hexachlorobutadiene	10
Hexachlorocyclopentadiene	10
Hexachloroethane	10
Hexachlorophen	--

Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270^a

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Parameter	Target Water Quantitation Limit (µg/l)
Hexachloropropene	20
Indeno(1,2,3-c,d)pyrene	10
Isophorone	10
Isosafrole	20
bis(2-chloroethoxy)Methane	10
Methapyrilene	10
3-Methylcholanthrene	20
Methyl methacrylate	10
Methyl methanesulfonate	10
2-Methylnaphthalene	10
Methyl parathion	--
Naphthalene	10
1,4-Naphthoquinone	10
1-Naphthylamine	10
2-Naphthylamine	10
2-Nitroaniline	50
3-Nitroaniline	50
4-Nitroaniline	50
Nitrobenzene	10
2-Nitrophenol	10
4-Nitrophenol	50

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Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270^a

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Parameter	Target Water Quantitation Limit (µg/l)
4-Nitroquinoline-1-oxide	--
N-Nitrosodi-n-butylamine	01
N-Nitrosodiethylamine	10
N-Nitrosodimethylamine	01
N-Nitrosodiphenylamine	10
N-Nitroso-di-n-propylamine	10
N-Nitrosomethylethylamine	10
N-Nitrosomorpholine	10
N-Nitrosopiperidine	10
N-Nitrosopyrrolidine	10
5-Nitro-o-toluidine	10
Pentachlorobenzene	10
Pentachloroethane	10
Pentachloronitrobenzene	50
Pentachlorophenol	50
Phenacetin	10
Phenanthrene	10
Phenol	10
p-Phenylenediamine	--
Phorate (Thimet)	--
bis(2-ethylhexyl)Phthalate	10

Table C.2
40 CFR 264 Appendix IX Semivolatile Organics
Method 8270^a

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Parameter	Target Water Quantitation Limit (µg/l)
2-Picoline	10
Pronamide	20
Pyrene	10
Pyridine	--
Safrole	10
Sulfotepp	--
1,2,4,5-Tetrachlorobenzene	10
2,3,4,6-Tetrachlorophenol	20
Thionazin	--
o-Toluidine	10
2,4,6-TGribromophenol	--
1,2,4-Trichlorobenzene	10
2,4,5-Trichlorophenol	50
2,4,6-Trichlorophenol	10
o,o,o-Triethylphosphorothioate	--
sym-Trinitrobenzene	10
^a Method 8270 must be modified to include all of the indicated parameters. Method 8270 is from EPA 1986.	

Table C.3
40 CFR 264 Appendix IX Organochlorine Pesticides/PCBs
Method 8080^a

Page 1 of 2

Parameter	Target Water Quantitation Limit (mg/l)
Aldrin	0.050
Aroclor-1016	0.50
Aroclor-1221	0.50
Aroclor-1232	0.50
Aroclor-1242	0.50
Aroclor-1248	0.50
Aroclor-1254	1.0
Aroclor-1260	1.0
alpha-BHC	0.050
beta-BHC	0.050
delta-BHC	0.050
gamma-BHC (Lindane)	0.050
Chlordane	0.50
Chlorobenzilate	0.50
4,4'-DDD	0.10
4,4'-DDE	0.10
4,4'-DDT	0.10
Diallate	1.5
Dieldrin	0.10

Table C.3
40 CFR 264 Appendix IX Organochlorine Pesticides/PCBs
Method 8080^a

Page 2 of 2

Parameter	Target Water Quantitation Limit (mg/l)
Endosulfan I	0.050
Endosulfan II	0.10
Endosulfan sulfate	0.10
Endrin	0.10
Endrin aldehyde	0.10
Heptachlor	0.050
Heptachlor epoxide	0.050
Isodrin	0.050
Kepone	0.10
Methoxychlor	0.50
Toxaphene	1.0
^a Method 8080 must be modified to include all of the listed parameters. Method 8080 is from EPA 1986.	

Table C.4
40 CFR 264 Appendix IX Herbicides
Method 8150^a

Parameter	Target Water Detection Limit (µg/l)
2,4-D(2,4-dichlorophenoxyacetic acid)	12
2,4,5-T (2,4,5-trichlorophenoxyacetic acid)	2
2,4,5-TP (Silvex; 2,4,5-trichlorophenoxypropanoic acid)	2
^a Method 8150 is from EPA 1986.	

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Table C.5
40 CFR 264 Appendix IX Dioxins and Furans
Method 8280^a

Parameters	Target Practical Quantitation Limits (ng/l)
Dioxins	
Total tetrachlorodibenzo-p-dioxins	10
2,3,7,8-tetrachlorodibenzo-p-dioxin	5
Total pentachlorodibenzo-p-dioxins	10
Total hexachlorodibenzo-p-dioxins	10
Total heptachlorodibenzo-p-dioxin ^b	--
Octachlorodibenzo-p-dioxin ^b	--
Furans	
Total tetrachlorodibenzofurans	10
Total pentachlorodibenzofurans	10
Total hexachlorodibenzofurans	10
Total heptachlorodibenzofuran ^b	--
Octachlorodibenzofuran ^b	--
^a Method 8280 is from EPA 1986. ^b These parameters are not on the 40 CFR 264 Appendix IX list, but have been added in accordance with the Liquid Effluent QAPP (WHC 1991).	

Table C.6
Metals

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Parameter	Target Water Detection Limit (mg/l)	Analytical Method ^a
Appendix IX Metals		
Antimony	0.032	6010
Arsenic	0.001	7060
Barium	0.002	6010
Beryllium	0.003	6010
Cadmium	0.004	6010
Chromium	0.007	6010
Cobalt	0.007	6010
Copper	0.006	6010
Lead	0.001	7421, 7420
Mercury	0.0002	7470
Nickel	0.015	6010
Selenium	0.002	7740
Silver	0.007	6010
Thallium	0.001	7841
Tin	0.8	7870
Vanadium	0.008	6010
Zinc	0.002	6010
Additional Metals		
Aluminum	0.045	6010
Calcium	0.01	6010
Iron	0.03	7380

Table C.6
Metals

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Parameter	Target Water Detection Limit (mg/l)	Analytical Method ^a
Magnesium	0.001	6010
Manganese	0.002	7460
Potassium	0.01	7610
Sodium	0.002	7770

^aMethods are from EPA 1986.

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Table C.7
General Parameters

Parameter	Target Water Detection Limit (mg/l)	Analytical Method
Appendix IX Parameters		
Cyanide, Total	0.01	335 ^a
Sulfide, Total	0.05	376 ^a
Additional Parameters		
Alkalinity	1.0	310 ^a
Ammonia	0.01	350 ^a
Chloride	1.0	325 ^a
COD (chemical oxygen demand)	9	410 ^a
Coliform (fecal)	1 colony/100 ml.	908C ^c
Coliform (total)	1 colony/100 ml.	908A ^c
Conductivity	--	9050 ^b
Cyanide (amenable)	0.01	335 ^a
Fluoride	0.1	340 ^a
Nitrate	0.03	352 ^a
Nitrite	0.02	354 ^a
pH	--	9045 ^b
Phenols (total)	--	420 ^a
Sulfate	1.0	375 ^a
Sulfite		377 ^a
TOC (total organic carbon)	1.0	9060 ^b
TOX (total organic halogens)	0.1	9020 ^b
Turbidity	--	180 ^a
^a Methods are from EPA 1983. ^b Methods are from EPA 1986. ^c Methods are from APHA 1989.		

APPENDIX C REFERENCES

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